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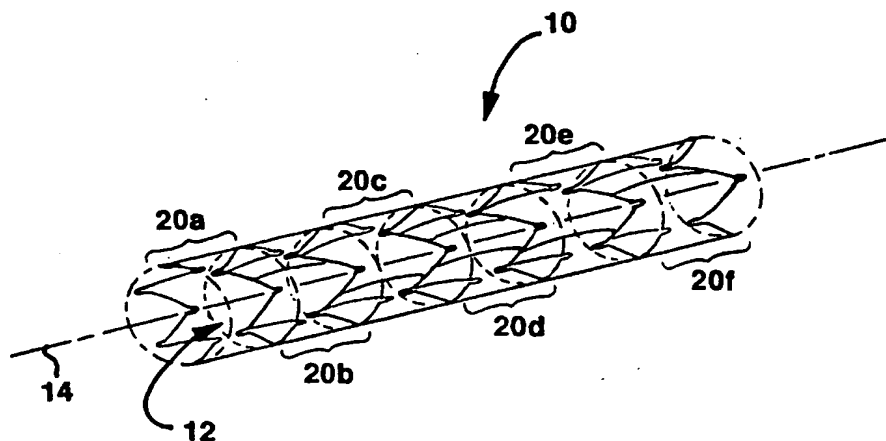
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(54) Title: **RADIALLY-EXPANDABLE STENT AND DELIVERY SYSTEM**

(57) Abstract

The present invention provides radially-expandable stents that, in various embodiments, may reduce the bending stresses/strains associated with the compressed state of self-expanding stents and/or may prevent longitudinal expansion/contraction of radially expandable stents between the compressed and expanded states. In addition, stents according to the present invention preferably exhibit increased longitudinal flexibility in both the compressed and expanded states. Furthermore, stents that include circumferential support sections with varying longitudinal lengths and/or varying unrestrained expanded diameters are also disclosed. Also disclosed are stents that provide controllable expansive force profiles. The present invention also includes delivery systems in which threading of the guidewire through the delivery system may be simplified. In addition, the delivery systems according to the present invention may also incorporate a balloon to assist in radially expanding the stent and/or seating of the stent in the lumen during deployment without removing the stent delivery catheter. Further, the delivery systems may also include a support tube at the proximal end to assist in fixing the position of the stent relative to a guide catheter during deployment of the stent.

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RADIALLY-EXPANDABLE STENT AND DELIVERY SYSTEM

Field of the Invention

5 The present invention relates to intravascular stent implants for maintaining vascular patency in humans and animals. More particularly, the present invention provides radially-expandable stents, delivery systems for deploying the stents within body lumens, and methods of deploying the stents.

Background of the Invention

10 Percutaneous transluminal coronary angioplasty (PTCA) is used to increase the lumen diameter of a coronary artery partially or totally obstructed by a build-up of cholesterol fats or atherosclerotic plaque. Typically a first guidewire of about .038 inches in diameter is steered through the vascular system to the site of therapy. A guiding catheter, for example, can then be advanced
15 over the first guidewire to a point just proximal of the stenosis. The first guidewire is then removed. A balloon catheter on a smaller .014 inch diameter second guidewire is advanced within the guiding catheter to a point just proximal of the stenosis. The second guidewire is advanced into the stenosis, followed by the balloon on the distal end of the catheter. The balloon is inflated causing the
20 site of the stenosis to widen.

The dilatation of the occlusion, however, can form flaps, fissures and dissections which threaten reclosure of the dilated vessel or even perforations in the vessel wall. Implantation of a stent can provide support for such flaps and dissections and thereby prevent reclosure of the vessel or provide a patch repair
25 for a perforated vessel wall until corrective surgery can be performed. It has also been shown that the use of intravascular stents can measurably decrease the incidence of restenosis after angioplasty thereby reducing the likelihood that a secondary angioplasty procedure or a surgical bypass operation will be necessary.

30 An implanted prosthesis such as a stent can preclude additional procedures and maintain vascular patency by mechanically supporting dilated vessels to prevent vessel reclosure. Stents can also be used to repair aneurysms,

to support artificial vessels as liners of vessels or to repair dissections. Stents are suited to the treatment of any body lumen, including the vas deferens, ducts of the gallbladder, prostate gland, trachea, bronchus and liver. The body lumens range in diameter from small coronary vessels of 3 mm or less to 28 mm in the aortic vessel. The invention applies to acute and chronic closure or reclosure of body lumens.

A typical stent is a cylindrically shaped wire formed device intended to act as a permanent prosthesis. A typical stent ranges from 5 mm to 50 mm in length. A stent is deployed in a body lumen from a radially compressed configuration into a radially expanded configuration which allows it to contact and support a body lumen. The stent can be made to be radially self-expanding or expandable by the use of an expansion device. The self expanding stent is made from a resilient springy material while the device expandable stent is made from a material which is plastically deformable. A plastically deformable stent can be implanted during a single angioplasty procedure by using a balloon catheter bearing a stent which has been crimped onto the balloon. The stent expands radially as the balloon is inflated, forcing the stent into contact with the interior of the body lumen thereby forming a supporting relationship with the vessel walls.

Conventional angioplasty balloons fall into high, medium and low pressure ranges. Low pressure balloons are those which fall into rated burst pressures below 6 atmospheres. Medium pressure balloons are those which fall into rated burst pressures between 6 and 12 atmospheres. High pressure balloons are those which fall into rated burst pressures above 12 atmospheres. Burst pressure is determined by material selection, wall thickness and tensile strength.

The biocompatible metal stent props open blocked coronary arteries, keeping them from reclosing after balloon angioplasty. A balloon of appropriate size and pressure is first used to open the lesion. The process is repeated with a stent crimped on a second balloon. The second balloon may be a high pressure type of balloon, e.g., more than 12 atmospheres, to insure that the stent is fully deployed upon inflation. The stent is deployed when the balloon is inflated. The stent remains as a permanent scaffold after the balloon is withdrawn. A high

pressure balloon is preferable for stent deployment because the stent must be forced against the artery's interior wall so that it will fully expand thereby precluding the ends of the stent from hanging down into the channel encouraging the formation of thrombus.

5 Various shapes of stents are known in the art. U.S. Patent No. 4,649,922 (Wiktor) discloses a linearly expandable spring-like stent. U.S. Patent No. 4,886,062 (Wiktor) discloses a two-dimensional zigzag form, typically a sinusoidal form. U.S. Patent No. 4,969,458 (Wiktor) discloses a stent wire
10 coiled into a limited number of turns wound in one direction, then reversed and wound in the opposite direction with the same number of turns, then reversed again and so on until a desired length is obtained.

 Stents have limited ability to provide effective patching of perforated vessels due to the spacing between metal elements. U.S. Patent No. 4,878,906 (Lindeman et al.) describes an endoprosthesis made of a thin wall molded plastic
15 sleeve intended to be collapsed radially and delivered to a damaged area of a vessel where it is expanded to provide a sealed interface to the vessel on its outer peripheral ends. The endoprosthesis therefore provides a patch which prevents leakage of blood from a vessel wall. The endoprosthesis disclosed employs various molded-in ribs, struts and the like to adapt the device for particular
20 applications and to provide the desired degree of stiffness to form the sealed interface with the vessel wall. Such a stiff prosthesis, however, could not be expected to have the longitudinal flexibility needed to adapt to curved vessels.

 One problem with self-expanding stents is that the stents must be compressed into a small diameter for delivery to the site or portion of the body
25 lumen at which support is desired. It is preferable that the stents be compressed into as small of a diameter as possible (typically referred to as "profile") to assist in delivering the stent to the desired site. That compression can, in some cases cause localized areas of high bending stress/strain within the stent.

 As a result of the high bending stresses/strain, the minimum profile for the
30 self-expanding stents can be limited to prevent non-recoverable strain levels in the stent and, therefore, ensure full radial expansion of the stent when released from

the delivery system. The larger profile can limit the delivery and use of the stent to larger diameter lumens.

Alternatively, if a small delivery profile is desired, then the stent may be designed to achieve that profile which can often result in a larger window area and a reduction in the outward forces generated by the stent after expansion within the lumen. The larger window area and, therefore, inferior body lumen scaffolding reduces the effectiveness against recurring restenosis. The reduced outward forces may be problematic if the stent does not firmly engage the wall of the lumen.

One attempt at addressing the high bending stresses/strains in a self-expanding stent is described in U.S. Patent No. 4,830,003 (Wolff et al.) in which the stent is made of a series of generally straight wire segments welded together at their ends to form a zigzag shaped stent when expanded. By using generally straight wires, the bending stresses/strains associated with bends in an integral wire-formed stent body can be avoided. Disadvantages associated with this approach include, however, the cost of manufacturing the stents by welding. The welds also lower the allowable stress levels in the stent, thereby limiting its fatigue life and compression for delivery. Another disadvantage is that the length of the stent can change significantly from the compressed state to the expanded state, thereby making accurate placement of the stent at the desired location within a body lumen more difficult.

Another attempt at addressing the high bending stresses/strains includes manufacturing self-expanding stents from materials other than metals as described in, e.g., U.S. Patent No. 5,356,423 (Tihon et al.). The stents disclosed therein are formed of thermoplastic materials and can be molded or otherwise formed into a fenestrated pattern similar to those produced by braided wire stents. By shaping the openings as depicted in Figure 5 of the patent, the stress concentration at the bending points can be reduced. Disadvantages of this approach include, however, degradation associated with implanted plastic materials, including changes in the elasticity of the plastics which can result in a reduction in the radially outward forces generated by the stent.

Another problem associated with self-expanding stents is that, particularly in connection with long lesions (e.g., those with a length of about 30 mm or more),

differences in vessel diameter between the proximal and distal ends of the lesion and/or stent may be greater than the desired operating range of a self-expanding stent. Furthermore, the increase in over-sizing from the proximal to the distal end of the stent may result in non-uniform pressure at the vessel wall along the length of the stent. That non-uniformity may cause insufficient or excessive pressure at one end of the stent after expansion.

Yet another problem with self-expanding stents is the need for stents that can provide varying outward force or pressure along their lengths. Attempts to provide stents with varying outward pressure profiles are typically limited to helically wound stents in which the coil density and/or helix angle are changed to affect the pressure generated by the stents after expansion. *See, e.g.,* U.S. Patent Nos. 5,246,445 (Yachia et al.) and 5,507,767 (Maeda et al.). These approaches are generally limited to gradual pressure changes and may be difficult to construct. In addition, changing the pitch of the helically wound stents can increase the size of the openings between adjacent windings, thereby reducing support of the vessel wall.

Another approach to providing a stent with a varying pressure profile is described in U.S. Patent No. 5,601,593 (Freitag) in which rows of Nitinol wires with different shape memories that allow for selective expansion of the different structures formed by the different wires. One problem with this approach is that the pressure generated between the activated wires can be significantly reduced, thereby reducing uniformity in the outward force along the entire length of the stent.

SUMMARY OF THE INVENTION

It is an object of the invention to provide a radially-expanding stent for implantation within a body lumen that provides for reductions in the bending stresses/strains associated with compression of the stent for delivery to the desired location within a body lumen.

It is another object of the present invention to provide a radially-expanding stent in which the longitudinal length of the stent remains unchanged from the compressed state to the expanded state.

It is a further object of the invention to provide a stent with improved longitudinal flexibility to allow for threading through tortuous lumens and lesions, as well as to permit implantation in highly curved lumens.

It is another object of the invention to provide a radially-expanding stent
5 that provides for controllable expansive force profile along the length of the stent.

It is another object of the invention to provide a radially-expanding stent with and unrestrained expanded diameter that varies along the length of the stent.

It is an object of some delivery systems according to the present invention to provide a delivery system in which the position of the stent can be fixed relative
10 to a guide catheter.

It is another object of some delivery systems according to the present invention to provide a balloon integral with the stent delivery device to allow for post-deployment dilatation of the stent without removing the stent delivery catheter.

15 It is another object of some delivery systems of the present invention to provide for simplified threading of a guidewire through a distal portion of a rapid-exchange delivery system.

In one aspect, the present invention provides radially expandable stent for implantation within a body lumen including an elongated generally tubular body
20 defining a passageway having a longitudinal axis; the body including a plurality of circumferential support sections arranged successively along the longitudinal axis, each of the support sections having a length along the longitudinal axis; each of the circumferential support sections including a plurality of primary bends interconnected by struts, the primary bends being located on alternating
25 ends of the support section around the circumference of the body, each of the struts connecting successive primary bends on opposite ends of the support section and having a midpoint generally located therebetween; and at least one longitudinal member connecting adjacent support sections in the body, the longitudinal member having a first end attached proximate the midpoint of one
30 of the struts and a second end attached proximate the midpoint of one of the struts in the adjacent support section;

wherein the stent is radially compressible into a compressed state in which the struts are generally aligned with the longitudinal axis and radially expandable into an expanded state in which the struts and the primary bends in each of the support sections are arranged in a zigzag pattern, and further wherein the

5 longitudinal length of the stent in the compressed state is substantially the same as the longitudinal length of the stent in the expanded state.

In another aspect the present invention provides a radially expandable stent for implantation within a body lumen including an elongated generally tubular body defining a passageway having a longitudinal axis, the body including at

10 least one circumferential support section having a length along the longitudinal axis; each of the circumferential support sections including a plurality of primary bends interconnected by struts, the primary bends being located on alternating ends of the support section around the circumference of the body, each of the struts connecting successive primary bends on opposite ends of the support

15 section and having a midpoint generally located therebetween; wherein the stent is radially compressible into a compressed state and radially expandable into an expanded state in which the struts and primary bends in each of the support sections are arranged in a zigzag pattern, and further wherein each pair of adjacent struts associated with each of the primary bends abut at a point between

20 the primary bend and the midpoint of each strut in the pair of adjacent struts when the stent is in the compressed state, whereby the bending stress is reduced at each primary bend of the plurality of primary bends.

In another aspect, the present invention provides a radially expandable stent for implantation within a body lumen including an elongated generally

25 tubular body defining a passageway having a longitudinal axis, the body including at least one circumferential support section having a length along the longitudinal axis; each of the circumferential support sections including a substantially continuous element including a plurality of primary bends interconnected by struts, the primary bends being located on alternating ends of

30 the support section around the circumference of the body, each of the struts connecting successive primary bends on opposite ends of the support section and having a midpoint generally located therebetween, wherein the stent is radially

compressible into a compressed state and radially expandable into an expanded state in which the struts and primary bends in each of the support sections are arranged in a zigzag pattern; and means for reducing bending stress at the primary bends when the stent is in the compressed state.

5 In another aspect, the present invention provides a delivery system for implantation of a radially-expandable stent within a body lumen including an inner tube having a proximal end and a distal end, the inner tube having an inner tube lumen formed therein, the inner tube lumen having an opening at the distal end of the inner tube; a cover sheath having a proximal end and a distal end, the
10 cover sheath comprising a wall defining a cover sheath lumen, the inner tube located within the cover sheath lumen; a stent positioned about the inner tube at the distal end of the cover sheath; a first guidewire opening in the inner tube lumen, the first guidewire opening spaced from the distal end of the inner tube; a second guidewire opening in the wall of the cover sheath, the second guidewire
15 opening located proximate the first guidewire opening; and a guide element having a distal end located within the inner tube lumen, the guide element extending between the first and second guidewire openings.

 In another aspect, the present invention provides a radially expandable stent for implantation within a body lumen including an elongated generally
20 tubular body defining a passageway having a longitudinal axis. The body includes a plurality of circumferential support sections arranged successively along the longitudinal axis, each of the support sections having a longitudinal length along the longitudinal axis. Each of the circumferential support sections includes a plurality of primary bends interconnected by struts, the primary bends
25 being located on alternating ends of the support section around the circumference of the body, each of the struts connecting successive primary bends on opposite ends of the support section. The stent further includes at least one longitudinal member connecting adjacent support sections in the body. Each of the circumferential support sections is radially compressible into a compressed
30 diameter and radially expandable into an unrestrained expanded diameter in which the struts and the primary bends in each of the support sections are

arranged in a zigzag pattern. In addition, the longitudinal lengths of at least two of the circumferential support sections are different.

In another aspect the present invention provides a radially expandable stent for implantation within a body lumen including an elongated generally tubular body defining a passageway having a longitudinal axis. The body includes a plurality of circumferential support sections arranged successively along the longitudinal axis, each of the support sections having a longitudinal length along the longitudinal axis. Each of the circumferential support sections includes a plurality of primary bends interconnected by struts, the primary bends being located on alternating ends of the support section around the circumference of the body, each of the struts connecting successive primary bends on opposite ends of the support section. The stent further includes at least one longitudinal member connecting adjacent support sections in the body. Each of the circumferential support sections is radially compressible into a compressed diameter and radially expandable into an unrestrained expanded diameter in which the struts and the primary bends in each of the support sections are arranged in a zigzag pattern, and further wherein the unrestrained expanded diameters of at least two of the support sections vary.

In another aspect, the present invention provides a radially expandable stent for implantation within a body lumen including an elongated generally tubular body defining a passageway having a longitudinal axis. The body includes a plurality of circumferential support sections arranged successively along the longitudinal axis, each of the support sections having a longitudinal length along the longitudinal axis. Each of the circumferential support sections includes a plurality of primary bends interconnected by struts, the primary bends being located on alternating ends of the support section around the circumference of the body, each of the struts connecting successive primary bends on opposite ends of the support section. The stent further includes at least one longitudinal member connecting adjacent support sections in the body. Each of the circumferential support sections is radially compressible into a compressed diameter and radially expandable into an unrestrained expanded diameter in which the struts and the primary bends in each of the support sections are

arranged in a zigzag pattern. In addition, the circumferential widths of the struts in at least two of the circumferential support sections are different.

In another aspect, the present invention provides a method of deploying a stent within a body lumen by providing a radially expandable stent including an elongated generally tubular body defining a passageway having a longitudinal axis. The body includes a plurality of circumferential support sections arranged successively along the longitudinal axis, each of the support sections having a longitudinal length along the longitudinal axis. Each of the circumferential support sections includes a plurality of primary bends interconnected by struts, the primary bends being located on alternating ends of the support section around the circumference of the body, each of the struts connecting successive primary bends on opposite ends of the support section. The stent further includes at least one longitudinal member connecting adjacent support sections in the body. Each of the circumferential support sections is radially compressible into the compressed diameter and radially expandable into an unrestrained expanded diameter in which the struts and the primary bends in each of the support sections are arranged in a zigzag pattern. In addition, the longitudinal lengths of at least two of the circumferential support sections are different. The method further includes advancing the distal end of the delivery device and the stent through a body lumen and deploying the stent at a desired location within the body lumen.

In another aspect, the present invention provides a method of deploying a stent within a body lumen by providing a radially expandable stent including an elongated generally tubular body defining a passageway having a longitudinal axis. The body includes a plurality of circumferential support sections arranged successively along the longitudinal axis, each of the support sections having a longitudinal length along the longitudinal axis. Each of the circumferential support sections includes a plurality of primary bends interconnected by struts, the primary bends being located on alternating ends of the support section around the circumference of the body, each of the struts connecting successive primary bends on opposite ends of the support section. The stent further includes at least one longitudinal member connecting adjacent support sections in the body. Each

of the circumferential support sections is radially compressible into the compressed diameter and radially expandable into an unrestrained expanded diameter in which the struts and the primary bends in each of the support sections are arranged in a zigzag pattern, and further wherein the unrestrained
5 expanded diameters of at least two of the support sections vary. The method further includes advancing the distal end of the delivery device and the stent through a body lumen and deploying the stent at a desired location within the body lumen.

In another aspect, the present invention provides a method of deploying a
10 stent within a body lumen by providing a radially expandable stent including an elongated generally tubular body defining a passageway having a longitudinal axis. The body includes a plurality of circumferential support sections arranged successively along the longitudinal axis, each of the support sections having a longitudinal length along the longitudinal axis. Each of the circumferential
15 support sections includes a plurality of primary bends interconnected by struts, the primary bends being located on alternating ends of the support section around the circumference of the body, each of the struts connecting successive primary bends on opposite ends of the support section. The stent further includes at least one longitudinal member connecting adjacent support sections in the body. Each
20 of the circumferential support sections is radially compressible into the compressed diameter and radially expandable into an unrestrained expanded diameter in which the struts and the primary bends in each of the support sections are arranged in a zigzag pattern. In addition, the circumferential widths of the struts in at least two of the circumferential support sections are different.
25 The method further includes advancing the distal end of the delivery device and the stent through a body lumen and deploying the stent at a desired location within the body lumen.

In another aspect, the present invention provides a method of deploying a
stent within a body lumen by providing a radially expandable stent on a delivery
30 system including an inner tube having a proximal end and a distal end, the inner tube having an inner tube lumen formed therein, the inner tube lumen having an opening at the distal end of the inner tube and a first guidewire opening in the

inner tube lumen, the first guidewire opening spaced from the distal end of the inner tube; a stent positioned on the exterior surface of the inner tube at the distal end of the inner tube; a cover sheath having a proximal end and a distal end, the cover sheath comprising a wall defining a cover sheath lumen, the inner tube and
5 stent located within the cover sheath lumen, the cover sheath further including a second guidewire opening in the wall of the cover sheath, the second guidewire opening located proximate the first guidewire opening in the inner tube; and a guide element having a distal end located within the inner tube lumen, the guide element extending between the first and second guidewire openings, wherein the
10 guide element comprises a guide lumen formed in the distal end of the guide element; positioning a guidewire within a body lumen, wherein a proximal end of the guidewire extends out of the body lumen; inserting the proximal end of the guidewire into the inner tube lumen at the distal end of the inner tube; advancing the proximal end of the guidewire through the inner tube lumen towards the first
15 guidewire opening and the distal end of the guide element, wherein at least a portion of the proximal end of the guidewire is advanced into the guide lumen in the distal end of the guide element; advancing the proximal end of the guidewire through the first and second guidewire openings; advancing the distal end of the inner tube and the stent over the guidewire towards the distal end of the
20 guidewire, wherein the stent is positioned at a desired location within the body lumen; and deploying the stent at the desired location within the body lumen.

In another aspect, the present invention provides a method of deploying a stent within a body lumen by providing a radially expandable stent on a delivery system including an inner tube having a proximal end and a distal end, the inner
25 tube having an inner tube lumen formed therein; a stent positioned on the exterior surface of the inner tube at the distal end of the inner tube; an expandable balloon located on the inner tube; an inflation lumen in fluid communication with the balloon, the inflation lumen extending from the proximal end of the delivery system to the balloon; and a cover sheath having a
30 proximal end and a distal end, the cover sheath comprising a wall defining a cover sheath lumen, the inner tube, stent, and balloon located within the cover sheath lumen; positioning the inner tube, stent, balloon and cover sheath within a

body lumen; moving the cover sheath proximally relative to the distal end of the inner tube to deploy the stent with the body lumen; and inflating the balloon within the stent.

In another aspect, the present invention provides a method of deploying a stent within a body lumen by providing a radially expandable stent on a delivery system including an inner tube having a proximal end and a distal end; a stent positioned on the exterior surface of the inner tube at the distal end of the inner tube; a cover sheath having a proximal end and a distal end, the cover sheath including a cover sheath lumen, the inner tube and stent located within the cover sheath lumen; and a support tube having a proximal end and a distal end, the support tube including a support tube lumen containing at least a portion of the proximal end of the cover sheath, the cover sheath being movable in the proximal and distal directions within the support tube lumen and the position of the inner tube being fixed relative to the position of the support tube; positioning a guide catheter within a body lumen; advancing the distal ends of the inner tube and the cover sheath through the guide catheter; fixing the position of the support tube relative to the guide catheter, wherein the positions of the distal end of the inner tube and the stent within the body lumen are also fixed relative to the guide catheter; and moving the cover sheath proximally to release the stent on the distal end of the inner tube, thereby deploying the stent within the body lumen.

These and other features and advantages of the present invention are described below in connection the description of the preferred embodiments.

BRIEF DESCRIPTION OF THE DRAWINGS

FIG. 1 is a perspective view of one radially expanded stent according to the present invention.

FIG. 2 is a plan view of the stent of Figure 1 in which the body of the stent is unrolled.

FIG. 3 is an enlarged partial view of the stent body of Figure 2 in the expanded state.

FIG. 4 is an enlarged partial view of the stent body of Figure 2 in the compressed state.

FIGS. 5-8 are enlarged partial views of portions of alternative stents according to the present invention.

FIG. 9 is a plan view of another stent according to the present invention in which the body of the stent is unrolled.

5 FIG. 10 is a plan view of another stent according to the present invention in which the body of the stent is unrolled.

FIGS 11A-11D illustrate different force profiles that can be provided by stents manufactured according to the present invention.

FIGS. 12 and 12A depict one method of manufacturing a conical stent.

10 FIGS. 12B-12D depict mandrels that may be used to manufacture stents having corresponding unrestrained expanded diameters.

FIG. 13 is a schematic diagram of one delivery system according to the present invention.

15 FIG. 13A is an enlarged diagram of one portion of the delivery system of Figure 13.

FIG. 13B is a cross-sectional view of the mechanism of Figure 13A taken along line 13B-13B in Figure 13A.

FIG. 14 is an enlarged cross-sectional view of the delivery system of Figure 13 taken along line 14-14 in Figure 13.

20 FIG. 15 is an enlarged cross-sectional view of the distal end of the delivery system of Figure 13.

FIG. 16 is an enlarged cross-sectional view of the distal end of an alternate delivery system incorporating a balloon.

25 FIG. 17 is an enlarged partial cross-sectional view of one rapid-exchange delivery system according to the present invention.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

The present invention provides stents that may reduce the bending stresses/strains associated with the compressed state of radially-expanding stents and/or may prevent longitudinal expansion/contraction of radially expandable stents between the compressed and expanded states. In addition, stents according

30

to the present invention preferably exhibit increased longitudinal flexibility in both the compressed and expanded states.

The stents according to the present invention may, in various embodiments, provide controllable expansive force profiles. The expansive force profiles can be controlled by varying the unrestrained expanded diameter of the stents, by varying the longitudinal length of the circumferential support sections in the stents, and/or by varying the circumferential width of the struts in the support sections.

The present invention also includes delivery systems in which threading of the guidewire through the delivery system may be simplified. In addition, the delivery systems according to the present invention may also incorporate a balloon to assist in radially expanding the stent and/or seating of the stent in the lumen during deployment without removing the stent delivery catheter. Further, the delivery systems may also include a support tube at the proximal end to assist in fixing the position of the stent relative to a guide catheter during deployment of the stent.

Although the following discussion, along with the figures, describes illustrative preferred embodiments and methods according to the present invention, those skilled in the art will understand that other structures and/or methods could also be used to accomplish the desired functions. For example, although stents having one or more support sections are described herein, it will be understood that stents manufactured according to the present invention could have any number of desired support sections needed to obtain a stent with a desired longitudinal length. Furthermore, it will be understood that the figures are schematic only, and that the relative dimensions of the various illustrated features are not intended to limit the scope of the present invention.

Figure 1 depicts one illustrative self-expanding stent according to the present invention. The depicted stent includes a generally tubular body 10 defining a passageway 12 extending along a longitudinal axis 14. The body 10 is preferably formed from a plurality of support sections 20a, 20b, 20c, 20d, 20e, and 20f (collectively referred to as support sections 20 below) arranged successively along the longitudinal axis 14. The body 10 is depicted in Figure 1 in its expanded state

in which the support sections 20 have been expanded radially outward from the longitudinal axis 14.

Figure 2 is a plan view of a portion of the body 10 of the stent depicted in Figure 1 in which the body has been unrolled from the tubular shape of Figure 1.

5 Each of the support sections 20 is depicted and has a length along the longitudinal axis 14. Referring specifically to support section 20a, the support section 20a includes a plurality of primary bends 22 and 22' located on alternating ends of the support section 20a. Primary bend 22 on one end of the support section 20a is connected to a primary bend 22' by a strut 24. Because of the alternating nature of
10 the primary bends 22 and 22', the primary bends 22/22' and struts 24 are arranged in a zigzag pattern when the stent is in the expanded state (as in Figures 1 and 2).

Adjacent support sections 20a and 20b are connected to each other by at least one longitudinal member 40 extending between the support sections 20a and 20b. It is preferred that the longitudinal members 40 are attached to the struts 24,
15 although they may be attached at any location on each of the support sections 20. More preferably, the longitudinal members 40 are attached to the struts 24 at the midpoint of the length of the struts 24 between the primary bends 22 and 22'. By attaching the longitudinal members 40 at the midpoints of the struts, the length of the body 10 of the stent along the longitudinal axis 14 will exhibit substantially no
20 change between the compressed state and the expanded state.

Although most of the adjacent support sections 20 are connected by only one longitudinal member 40 in Figure 2, it should be noted that a plurality of longitudinal members 40 can be used to connect the support sections 20. For example, support sections 20c and 20d are connected by two longitudinal members
25 40 in Figure 2. Where more than one longitudinal member 40 is used to connect adjacent support sections 20, it is preferred that the longitudinal members be spaced evenly about the circumference of the body 10. For example, where two longitudinal members 40 are provided, it is preferred that they be located about 180 degrees apart, three longitudinal members 40 would preferably be located about
30 120 degrees apart, etc.

It may be preferred (but not required) that smaller stents, i.e., those having a diameter as manufactured of about 6 millimeters or less, employ two or more

longitudinal members 40 to connect adjacent support sections 20. It may also be preferred (but not required) that larger stents, i.e., those having manufactured diameters of about 5 millimeters or more, employ three or more longitudinal members 40 to connect adjacent support sections 20. It may further be preferred
5 that the number of longitudinal members 40 be less than the number of primary bends 22 found in each of the adjacent support sections 20. By reducing the number of longitudinal members 40, the longitudinal flexibility of the stents can be improved. The exact number of longitudinal members used in any stent according to the present invention will, however, vary based on the need for longitudinal
10 flexibility

It is preferred that the longitudinal members 40 connecting immediately adjacent support sections 20 are not aligned along the longitudinal axis 14 of the stent. As one example of this, the arrangement of the longitudinal members 40 in the first three support sections 20a, 20b, and 20c can be described. As can be seen
15 in Figure 2, the longitudinal members 40 connecting support sections 20a and 20b are not aligned along the longitudinal axis 14 with the longitudinal members 40 connecting support sections 20b and 20c. It is generally preferred that the longitudinal members connecting, e.g., support sections 20a and 20b, be tangentially out of phase from longitudinal members 40 connecting support
20 sections 20b and 20c by as large an amount as possible. For example, if the support sections 20a, 20b and 20c are all connected to their adjacent support sections 20 by two longitudinal members 180 degrees apart, it is preferred that the longitudinal members 40 connecting support sections 20a and 20b be tangentially out of phase by 90 degrees from the longitudinal members 40 connecting the
25 support sections 20b and 20c.

Providing longitudinal members 40 connecting immediately adjacent support sections 20, e.g., 20a and 20b, circumferentially spaced about the support sections 20 can improve the flexibility of the body 10 along the longitudinal axis 14. In addition, providing the longitudinal members 40 tangentially out of phase
30 along the length of the body 10, e.g., between sections 20a-20b and 20b-20c, can also improve the longitudinal flexibility of stents according to the present invention. Although these concepts have been described with reference to three

successive support sections, it will be understood that these concepts can be extended along the entire length of a stent incorporating as few as two support sections and as many support sections as desired.

It is significant to note that the longitudinal bending flexibility is improved both when the stent is in the compressed state during delivery and upon deployment of the stent in a body lumen. Increased longitudinal bending flexibility when compressed permits threading of the stent through long tortuous vessels and lesions. Increased longitudinal bending flexibility when expanded allows for deployment in highly curved vessels or lumens.

It will be understood that the longitudinal members 40 described above may be incorporated into self-expanding stents or into stents that are not self-expanding, i.e., stents that must be expanded by a balloon or some other method. In addition, the connection of the longitudinal members 40 can be used in any stent providing zigzag support sections, whether the stent includes primary bends such as those described herein or not. In any case, the connection of the longitudinal members 40 the midpoints of the struts 24 in adjacent zigzag support sections 20 will prevent changes in the longitudinal length of stents incorporating zigzag support sections similar to those described herein.

Figure 3 is an enlarged view of a portion of the one of the support sections 20 in Figure 2. A primary bend 22 is shown along with two opposing primary bends 22' on the opposite end of the support section. The primary bend 22 is attached to a pair of struts 24a and 24b. The lower strut 24a is attached to the lower opposing primary bend 22' while the upper strut 24b is attached to the upper opposing primary bend 22'. Strut 24a has a midpoint 25a that is generally located midway between the primary bend 22 and the lower opposing primary bend 22' while strut 24b has a midpoint 25b that is generally located midway between the primary bend 22 and the upper opposing primary bend 22'.

Strut 24a includes a secondary bend 26a located between its midpoint 25a and the primary bend 22. The secondary bend 26a forms an apex 27a facing the other strut 24b attached to the primary bend 22. Strut 24b includes a secondary bend 26b located between its midpoint 25b and the primary bend 22. The secondary bend 26b forms an apex 27b facing the other strut 24a attached to the

primary bend 22. As depicted in Figures 2 and 3, it is preferred that each of the struts 24 connecting primary bends 22 and 22' include two secondary bends, with one secondary bend being located on each side of the midpoint of the strut 24.

Figure 4 depicts the portion of the support section of Figure 3 in the compressed state in which the opposing upper and lower primary bends 22' are moved together. As a result, the struts 24a and 24b are also moved together, and abut each other first at a point between the midpoints 25a and 25b of the respective struts 24a and 24b. In the embodiment depicted in Figures 3 and 4, the point at which the struts associated with or attached to the primary bend 22 abut first is at the apexes 27a and 27b of the struts 24a and 24b. As a result, the minimum radius formed by the primary bend 22 during compression of the stent is limited by the abutting relationship of the apexes 27a and 27b that redistributes the stresses associated with compression of the stent into the struts 24a and 24b.

The construction of the supports sections depicted in Figures 2-4 can be modified while still limiting the maximum stresses associated with compression of the stent. One alternative is depicted in Figure 5 and includes a primary bend 122 on one end of a support section and two opposing primary bends 122' on the opposing end of the support section. The strut 124a connecting the primary bend 122 with the lower opposing primary bend 122' includes a secondary bend 126a and the strut 124b connecting the primary bend 122 with the upper opposing primary bend 122' includes a secondary bend 126b.

The primary difference between the embodiments depicted in Figures 3 and 5 is that the portion 123a of the strut 124a between the secondary bend 126a and the primary bend 122 is not generally parallel to the corresponding portion 123b of the strut 124b. As described with respect to the embodiment of Figure 3 above, however, it is preferred that the struts 124a and 124b abut first at the apexes 127a and 127b formed by the secondary bends 126a and 126b during compression to thereby reduce the maximum bending stresses associated with compression of the stent.

Another alternative construction is depicted in Figure 6 and includes a primary bend 222 on one end of a support section and two opposing primary bends 222' on the opposing end of the support section. The strut 224a connecting the

primary bend 222 with the lower opposing primary bend 222' includes a secondary bend 226a and the strut 224b connecting the primary bend 222 with the upper opposing primary bend 222' includes a secondary bend 226b.

5 The primary bend 222 in the embodiment of Figure 6 and the portions of the struts 224a and 224b located between the secondary bends 126a and 126b and the primary bend 222 form a generally circular element as seen in Figure 6. As described with respect to the embodiments of Figures 3 and 5 above, however, it is preferred that the struts 224a and 224b abut first at the apexes 227a and 227b formed by the secondary bends 226a and 226b during compression to thereby
10 reduce the maximum bending stresses associated with compression of the stent.

Yet another alternative construction is depicted in Figure 7 and includes a primary bend 322 on one end of a support section and two opposing primary bends 322' on the opposing end of the support section. The strut 324a connecting the primary bend 322 with the lower opposing primary bend 322' includes a protrusion
15 327a facing the opposing strut 324b attached to the primary bend 322. Similarly, the strut 324b connecting the primary bend 322 with the upper opposing primary bend 322' includes a protrusion 327b facing the opposing strut 324a.

Although the struts 324a and 324b do not include secondary bends as in those struts described above, the protrusions 327a and 327b define the point at
20 which the struts 324a/324b first abut when the stent is compressed. Because that point is removed from the primary bend 322, the minimum bending radius of the primary bend is limited, thereby reducing the maximum bending stresses at the primary bends that is associated with compression of the stent.

Figure 8 illustrates yet another feature of stents according to the present
25 invention when compared to the stent depicted in Figure 2. The view of Figure 8 is a portion of a stent body including two adjacent support sections 420a and 420b. Support section 420a includes primary bends 422a on one end and opposing primary bends 422a' on the opposite end of the support section 420a. Similarly, the support section 420b includes primary bends 422b on one end of the support
30 section 420b and opposing primary bends 422b' on the opposite end of the support section 420b.

In the embodiment depicted in Figure 8, the primary bends 422a and 422b in adjacent support sections 420a and 420b are generally aligned along the longitudinal axis 414. Likewise, the primary bends 422a' and 422b' in adjacent support sections 420a and 420b are also generally aligned along the longitudinal axis 414. As a result, the support sections 420a and 420b are said to be "in phase" with each other.

Figure 2 illustrates a stent in which the support sections 20 are "out of phase" with the adjacent support sections because the primary bends 22 and 22' on the adjacent support sections 20 do not generally align along the longitudinal axis 14 as do the primary bends 422a/422b and 422a'/422b' in the embodiment depicted in Figure 8.

Figure 9 illustrates another variation that may be advantageous in connection with stents according to the present invention. Figure 9 is a plan view in which the body of a stent 510 has been unrolled from its tubular shape (after expansion). Each of the support sections 520 has a length along the longitudinal axis 514. The support sections 520a-520n (referred to generally as support sections 520) are connected to adjacent support sections along the longitudinal axis 514 by at least one longitudinal member 540 extending between pairs of adjacent support sections 520.

The stent 510 is able to compensate for variations in vessel diameter over its length by increasing the longitudinal length of the support sections 520 over the length of the stent 520. In the illustrated stent 510, the longitudinal length of each adjacent support section 520 increases from section 520a to section 520n. The support sections 520 with different longitudinal lengths exert different outward forces against the vessel wall after expansion. As a result, the differing lengths of the support sections 520 increases the permissible over-sizing of the stent 510 within a vessel. In other words, the stent 510 can be implanted into a successively smaller diameter vessel while limiting the amount of overstressing of the stent 510 after expansion. In addition, although the distal end of the stent 510 may be compressed to a greater degree after expansion in a narrowing vessel, the outward force or pressure applied to the vessel by the longer support sections 520 in the narrower distal portion of the vessel is smaller, thereby resulting in more uniform

applied pressure over the length of the stent 510 (as compared to a stent in which all of the support sections have the same longitudinal length).

The increases in longitudinal length between adjacent support sections 520 in the illustrated stent 510 are substantially constant over the entire stent 510. In another alternative, the differences in longitudinal length may increase along the length of the stent 510, i.e., the difference in length between support sections 520a and 520b is less than the difference between support sections 520b and 520c. Those variations in longitudinal length difference between adjacent support sections 520 can be uniform (e.g., linear, exponential, etc.) or they may be polynomial, random, etc. In another alternative, length of some sections 520 may be the same, with the length of only some support sections 520 varying.

Variations in the lengths of support sections in stents manufactured according to the present invention may be controlled to provide controlled variations in the outward forces provided by the stents along their lengths, whether the stents are located in vessels having varying or uniform diameters. For example, it may be desirable to provide a stent in which the outward forces generated at each end of the stent are reduced, thereby also providing a stent with reduced radial stiffness at its ends.

One such stent 610 is illustrated in Figure 10 in which the support sections 620a, 620b, 620m and 620n have an increased length along the longitudinal axis 614 as compared to the interior support sections 620c-620l. One benefit of this design is that the ends of the stent 610 may have a reduced longitudinal stiffness which may reduce injury to the vessel walls at the ends of the stent 610. That reduction in vessel injury may provide a decrease in restenosis rates in connection with the stent 610.

Figures 11A-11D illustrate a few different force profiles in which the horizontal axis is indicative of the longitudinal length of the stent and the vertical axis is indicative of the outward force or pressure exerted by the stent on a vessel wall after expansion. The depicted force profiles assume that the stents are uniformly constrained along their lengths after expansion within a vessel, regardless of whether the vessel has a constant or varying inner diameter.

The force profile illustrated in Figure 11A is indicative of the force profile to be expected from a stent similar to stent 510, i.e., a stent in which successive support sections 520 generate continually decreasing outward forces. The force profile illustrated in Figure 11B is indicative of the force profile to be expected
5 from a stent similar to stent 610, i.e., a stent in which the support section or sections at the ends of the stent generate reduced outward forces as compared to the central portion of the stent in which the outward forces are substantially uniform.

The force profile illustrated in Figure 11C is indicative of a stent that
10 combines the features of stent 510 with stent 610. As illustrated, the outward forces generated by the support section or sections at the ends of the stent are reduced as compared to the immediately adjacent support sections. In addition, however, the support sections in the central portion of the stent generate continually reducing outward forces moving along the length of the stent from left
15 to right in Figure 11C.

The force profile illustrated in Figure 11D is indicative of yet another variation in the outward force profiles possible using stents manufactured according to the present invention. As seen in Figure 11D, the outward forces generated by the support sections reach a peak near the center of the stent and
20 gradually decrease towards each end of the stent.

Figures 12 and 12A illustrate another variation in stents according to the present invention. The stent 710 is formed with an expanded shape that is substantially conical. In the preferred stent 710 the support sections 720a-720e have a substantially constant length along the longitudinal axis 714. If the stent
25 710 is manufactured from a material such as a nickel titanium alloy (e.g., Nitinol), the stent 710 is preferably manufactured at the smaller diameter and then placed on a conical mandrel 790 (see Figure 12) to increase its diameter. By proper heat treatment of the Nitinol during manufacturing, the conical shape can be imparted into the "memory" of the stent 710. Upon expansion in a vessel, one end of the
30 stent 710 will be urged to a larger diameter than the other end of the stent 710. As a result, the outward force or pressure exerted on a vessel wall with a narrowing diameter will preferably be more uniform with stent 710 than with a stent having a

uniform expanded diameter. Furthermore, stent 710 can be used to generate a force profile similar to that illustrated in Figure 11A when expanded in a vessel which does not have a diameter that narrows at the same rate at which the diameter of the stent 710 narrows, even though the support sections 720 of the stent are all
5 of substantially the same longitudinal length.

Yet another manner in which the outward expansive forces generated by the stents manufactured according to the present invention may vary is based on the circumferential width of each of the struts located between bends in each of the support sections. Each of the struts, e.g., strut 24 in Figure 2, has a width as
10 measured circumferentially around the formed stent. In addition, each of the struts 24 has a thickness measured radially from the longitudinal axis 14 (See Figure 1) of the formed stent.

By varying the circumferential width of the struts 24 both along their length and through each of the bends 22 and 22', some control over the expansive forces
15 generated by the support sections 20 may be achieved. For example, a support section 20 with wider struts 24 and bends 22/22' may provide a larger expansive force than an otherwise comparable support section in which the circumferential width of the struts is smaller. These variations in strut width can be provided between adjacent support sections 20 and the strut width may vary between support
20 sections along the stents to provide the desired expansive force profiles, examples of which are illustrated in Figures 11A-11D. Furthermore, variations in circumferential width of the support struts can be combined with variations in longitudinal lengths of the support sections and/or expanded diameter to yield desired expansive force profiles.

25 When stents are manufactured of a shape-memory metal, such as Nitinol, that also exhibits superelasticity, the shape of the mandrel upon which the stents are formed can be used to provide varying pressure profiles in a stent, whether the stent has support sections with different lengths or substantially uniform longitudinal lengths. The support sections of stents formed on the mandrels
30 illustrated in Figures 12A-12D will exhibit unrestrained expanded diameters based on the diameter of that portion of the mandrel on which they are formed. When employed in, e.g., a vessel having a generally uniform diameter, the support

sections formed with the different unrestrained expanded diameters illustrated by the mandrels of Figures 12A-12D will exhibit force profiles similar to their unrestrained expanded diameters. For example mandrel 790' in Figure 12B could be used to manufacture a stent that can generate a force profile similar to that illustrated in Figure 11B when expanded (i.e., constrained) in a vessel with a uniform diameter. Mandrel 790'' in Figure 12C could be used to form a stent that generated the force profile illustrated in Figure 11C when constrained within a vessel with a generally uniform diameter after expansion. Similarly, the mandrel 790''' in Figure 12D could be used to manufacture a stent exhibiting the force profile illustrated in Figure 11D when constrained within a vessel having a generally uniform diameter after expansion.

The radially expandable stents depicted and described above with respect to Figures 1-12 are preferably formed as a one-piece, completely integral units from a thin-walled tube of suitable material. Typically, the stents will be cut or machined from a tube using, e.g., laser, water jet, EDM (electrical discharge machining), or chemical etching techniques. As a result, the stents can be formed without welds or joints. It is also envisioned, however, that stents according to the present invention could be formed from a sheet of material using, e.g., laser, water jet, EDM, or chemical etching techniques. If the stent was formed from a sheet of material, the body 10 as seen in Figure 2 would be formed into a tube and welded or otherwise joined along one side of the stent resulting in a series of welds or other joints along the length of the body.

Preferred materials for stents according to the present invention include those materials that can provide the desired functional characteristics with respect to biological compatibility, modulus of elasticity, etc. For example, it is preferred that the stents be biologically compatible, as well as be capable of significant recoverable strain to assume a low profile for delivery to a desired location within a body lumen. After release of the compressed stent, it is preferred that the stent be capable of radially expanding back to its original diameter.

Particularly preferred materials for stents according to the present invention are nickel titanium alloys and other alloys that exhibit superelastic behavior, i.e., are capable of significant distortion without plastic deformation. Stents

manufactured of such materials can be significantly compressed without plastic deformation, i.e., they are compressed such that the maximum strain level in the stent is below the recoverable strain limit of the material. Discussions relating to nickel titanium alloys and other alloys that exhibit behaviors suitable for stents according to the present invention can be found in, e.g., U.S. Patent No. 5,597,378 (Jervis) and WO 95/31945 (Burmeister et al.). Nickel titanium alloys and tubes suitable for use in manufacturing stents according to the present invention can be obtained from, e.g., Memry Corp., Menlo Park, California; Minitubes, Grenoble, France; NDC, Fremont, California; and Shape Memory Applications, Inc., Santa Clara, California.

The radially outward directed force developed by the stents according to the present invention serves two functions. One function is to hold the body lumen open against a force directed radially inward, e.g., a spasm, as well as preventing restriction of the passageway through the lumen by intimal flaps or dissections generated by, e.g., prior balloon angioplasty. Another function is to fix the position of the stent within the body lumen by intimate contact between the stent and the walls of the lumen. The outwardly directed forces must not be excessive, however, to avoid traumatization of the lumen walls by the stent.

The diameters of some preferred stents when in the compressed state for delivery to a desired location within a body lumen is typically reduced from about two to about six times the diameter of the stents when in their expanded state before compression. For example, typical stents may have a compressed external diameter of about 1 millimeter to about 3 millimeters for delivery and an expanded external diameter in a body lumen of about 3 millimeters to about 15 millimeters when released from compression in a large arterial vessel. Some preferred stents used in coronary arteries may have a compressed external diameter of about 1 millimeter and an expanded external diameter in a body lumen of up to about 6 millimeters.

In addition to ranges in diameters, it will also be understood that the stents according to the present invention can have any desired longitudinal length as required for a particular application. Furthermore, although the illustrative stents depicted in Figures 1-12 have a plurality of successive support sections, it will be

understood that some stents according to the present invention could be manufactured with only one support section (in which case no longitudinal members would be required to connect adjacent support sections).

Having thus described radially expandable stents according to the present invention, we will now describe one delivery system suitable for deploying the self-expanding stents described above as well as other radially expandable stents. The delivery system depicted in Figures 13-15 provides for delivery of a stent to a desired location within a body lumen. It will be understood that the stents described above may be deployed by any suitable delivery system and they are not to be limited to deployment by the delivery systems described below.

The delivery system of Figure 13 includes a handle 50 at the proximal end. The handle 50 includes a release button 51 that slides within a channel 52 located in the handle 50. Preferably, the release button 51 is actuated by a user's thumb to assist in one-handed delivery of the stent as discussed in more detail below.

It is preferred that the release button 51 be locked or retained in position before delivery to prevent accidental or unwanted deployment of the stent from the delivery system. Referring to Figure 13A, one preferred retaining mechanism is a bend or turn in the distal end of the channel 52 such that the channel 52 includes a circumferential portion 52a at the distal end connecting to the otherwise longitudinal channel 52 seen in Figures 13 and 13A. Retaining the release button 51 in position at the distal end of the channel 52 (in the circumferential portion of the channel) until delivery of the stent is desired, at which time the button 51 is moved circumferentially and then longitudinally along the length of the channel 52 to release the stent as discussed in more detail below.

The illustrated channel 52 also includes two constrictions 53a and 53b that are also provided to assist in controlling movement of the release button 51. The constrictions 53a and 53b are narrowed such that an increased force is required to move the release button 51 past the localized narrowing. Constriction 53a works in combination with circumferential portion 52a of the channel to retain the button in position at the distal end of the channel 52 while constriction 53b assists in retaining the release button 51 near the proximal end of the channel 52 after deployment of a stent.

Those skilled in the art will understand that a variety of retaining mechanisms could be substituted for the preferred mechanism described above. Examples of suitable alternatives include, but are not limited to: a removable security band around the handle 50 that must be removed to move the release button 51 proximally, stoppers within the channel 52 that must be removed to move the release button 51 proximally, a detent mechanism in which the release button can be depressed radially inward to release the button 51 for movement within the channel, etc.

Figure 13B illustrates another feature of the deployment mechanism, i.e., the inclusion of an anti-reverse mechanism. By anti-reverse, it is meant that movement of the button 51 in the opposite direction (distally) after deployment has begun is restricted or prevented. The illustrated anti-reverse mechanism seen in Figure 13B includes a flat bent wire spring 48 that contacts the inner surface 49 of the handle. As the button 51 is moved proximally (to the right in Figure 13B), wire 48 drags along the surface 49. Attempts to move the button 51 distally (to the left in Figure 13B) will, however, cause the end of the wire 48 to become embedded in the inner surface 49, thereby restricting or preventing such movement. Although wire 48 provides one embodiment of an anti-reverse mechanism, those skilled in the art will recognize many other mechanisms that perform the same or substantially the same function including, but not limited to ratcheting mechanisms, silicone seals, etc.

A support tube 54 extends from the distal end of the handle 50 and preferably extends into the hemostasis valve 94 of the Y-connector 92 of a guide catheter 90. Preferably, the support tube 54 terminates within the guide catheter 90 at a point near the Y-connector 92. The guide catheter 90 preferably terminates at a distal end 96 spaced from the Y-connector 92. The construction of guide catheters, Y-connectors and hemostasis valves are well known and will not be described further.

Figure 14 is a cross-sectional view of the proximal portion of the delivery system taken along the longitudinal axis of the support tube 54 as indicated by line 14-14 in Figure 13. The support tube 54 is coaxial with a cover sheath 70 and inner tube 60, both of which are described in more detail below. It is preferred that

the cover sheath 70 be movable within the support tube 54 and that the cover sheath 70 also be movable relative to the inner tube 60. Further, it is preferred that the inner tube 60 and the support tube 54 be fixed relative to each other.

Figure 15 is an enlarged view of the distal portion of the delivery device in which the stent 10 is located within the lumen 72 formed by the cover sheath 70. The cover sheath 70 maintains the stent 10 in a compressed state in which the stent 10 has a diameter suitable for delivery to an internal body lumen 100. Because the stent 10 is self-expanding, it is biased radially outward against the interior surface of the cover sheath 70 as depicted.

10 An inner tube 60 preferably extends through the cover sheath 70 and the compressed stent 10 as seen in Figure 15. The inner tube 60 also preferably includes a guidewire lumen 64 extending through to the distal end 61 of the inner tube 60. For clarity, the guidewire 104 has been removed from the guidewire lumen 64 in the inner tube 60 of Figure 15.

15 The preferred inner tube 60 includes a shoulder 62 proximal to the proximal end 16 of the stent 10. The shoulder 62 prevents the stent 10 from moving proximally with the cover sheath 70 during deployment because the outside diameter of the inner tube 60 at the shoulder 62 is greater than the inside diameter of the compressed stent 10. As a result, the position of the stent 10 relative to the shoulder 62 on inner tube 60 is fixed when the cover sheath 70 is moved proximally during deployment of the stent 10 as described below.

20 Inner tube 60 preferably extends to the handle 50 of the delivery system depicted in Figure 13. Furthermore, the inner tube 60 is preferably fixedly attached to the handle 50 and is substantially inextensible along its length. As a result, the distance between the handle 50 and the shoulder 62 on the inner tube 60 is fixed. Because the distance between the shoulder 62 and the handle 50 is fixed, the distance between the handle 50 and the compressed stent 10 on the interior surface of the cover sheath 70 is also fixed during deployment.

25 The stent 10 is released by moving the cover sheath 70 towards the proximal end of the delivery device, i.e., away from the distal end 61 of the inner tube 60. The cover sheath 70 is connected to an actuator such as a release button 51 on the handle 50 such that movement of the button 51 towards the proximal end

53 of the handle 50 moves the cover sheath 70 in the proximal direction towards the handle 50. If the stent 10 is self-expanding, that movement of the cover sheath 70 preferably removes the constraining forces on the compressed stent 10, thereby allowing it to expand within the lumen 100. Actuators that accomplish the
5 function of moving the cover sheath 70 relative to the handle 50 other than the preferred release button 51 will be known to those skilled in the art.

To assist in positioning the stent 10 during delivery, it is preferred that one radio-opaque marker 68 be provided on the inner tube 60 at the proximal end 16 of the stent 10 and another radio-opaque marker 74 be provided on the cover sheath
10 70 at the distal end 18 of the stent 10. Movement of the marker 74 on the cover sheath 70 past the marker 68 on the inner tube 60 is preferably indicative of sufficient movement of the cover sheath 70 such that the stent 10 is no longer constrained by within the lumen 72 of the cover sheath has been deployed within the body lumen 100.

15 Figure 16 is an enlarged view of the distal portion of an alternative delivery system incorporating an inflatable balloon 180 on the inner tube 160, with the balloon preferably located within the passageway formed by the compressed stent 110. As described in connection with the embodiment depicted in Figure 15, the stent 110 is located within the lumen 172 formed by the cover sheath 170. The
20 cover sheath 170 maintains the stent 110 in a compressed state in which the stent 110 has a diameter suitable for delivery to an internal body lumen. Because the stent 110 is self-expanding, it is biased radially outward against the interior surface of the cover sheath as depicted.

The inner tube 160 also preferably includes a guidewire lumen 164
25 extending through to the distal end 161 of the inner tube 160. The inner tube 160 also includes a shoulder 162 at the proximal end 116 of the stent 110 to assist in deploying the stent 110 as described above in connection with Figure 15. Inner tube 160 also preferably extends to the handle of a delivery system as described above in connection with Figure 15.

30 As seen in Figure 16, the portion of the inner tube 160 on which the balloon 180 is mounted preferably has a reduced diameter to maintain a low profile while allowing room for the balloon 180. The inner tube 160 also includes an

inflation lumen 182 in fluid communication with the interior of the collapsed balloon 180. The inflation lumen 182 is used to deliver the fluids used to inflate the balloon 180 during deployment of the stent 110. The inflation lumen 182 preferably terminates at the proximal end of the inner tube 160 where the fluid
5 source can be connected by known methods.

To assist in positioning the stent 110 during delivery, it is preferred that one radio-opaque marker 168 be provided on the inner tube 160 at the proximal end 116 of the stent 110 and another radio-opaque marker 174 be provided on the cover sheath 170 at the distal end 118 of the stent 110. Movement of the marker
10 174 on the cover sheath 170 past the marker 168 on the inner tube 160 is preferably indicative of sufficient movement of the cover sheath 170 such that the stent 110 is no longer constrained by within the lumen 172 of the cover sheath has been deployed within a body lumen.

As described above in connection with Figure 13, the preferred delivery
15 systems according to the present invention also preferably include a support tube 54 exterior to and coaxial with the cover sheath 70 and inner tube 60 to further assist in accurate placement of the stent 10. The support tube 54 preferably extends from the handle 50 and is sufficiently long to extend into the lumen of the guide catheter 90. As best seen in Figure 13, the support tube 54 preferably extends into,
20 e.g., a Y-connector 92 of the guide catheter 90 such that the position of the support tube 54 can be fixed relative to the guide catheter 90 by closure of the hemostasis valve 94 on the Y-connector 92.

It is preferred that the support tube 54 be fixedly attached to the handle 50 and that the support tube 54 be substantially inextensible along its longitudinal axis
25 such that, after the support tube 54 is fixed in the hemostasis valve 94, the handle 50 is located a fixed distance from the hemostasis valve 94. The cover sheath 70 located within the support tube 54 (see Figure 14) is, however, free to move longitudinally within the support tube 54 during deployment of the stent 10. Because the support tube 54 and the inner tube 60 are both fixedly attached to the
30 handle 50, however, the distance between the stent 10 and the hemostasis valve 94 (and handle 50) are also fixed on closure of the hemostasis valve 94 on the support tube 54.

Use of the delivery system described above will now be described in connection with balloon angioplasty treatment of a lesion within a coronary vessel. Deployment of the stent will typically involve balloon angioplasty to expand the passageway through a lesion. Typically, a balloon catheter will be advanced over a
5 guidewire to the desired location. After dilatation, the balloon catheter will be withdrawn while the guidewire 104 and guide catheter 90 used with the balloon catheter remain in position. The guide catheter 90 is typically sutured in position to fix its location relative to the patient. At that point, the inner tube 60 and cover sheath 70 with compressed stent 10 will be advanced through the guide catheter 90
10 past the distal end 96 of the guide catheter 90 along the guidewire 104 until the stent 10 is in the desired location relative to the lesion 102. That position can be verified by, e.g., using the radio-opaque markers 68 and 74 on the inner tube 60 and cover sheath 70 as described above.

With the stent 10 in the desired location, the hemostasis valve 94 is
15 preferably fastened or closed on the support tube 54, thereby fixing the position of the stent relative to the guide catheter 90 (which, in turn fixes the position of the stent 10 relative to the patient because of the connection between the guide catheter 90 and the patient as described above). With the hemostasis valve 94 closed, the release button 51 is moved from its locked position within the channel 52 and then
20 gently moved towards the proximal end 53 of the handle 50. That movement preferably draws the distal end 76 of the cover sheath 70 past the stent 10. If the stent 10 is self-expanding, it will typically expand radially outward from the inner tube 60 towards the interior surface of the lesion 102.

After the cover sheath 70 is withdrawn sufficiently to expose the stent 10,
25 the balloon 80 can be inflated to either expand the stent 10 (if it is not self-expanding) or to assist in proper seating of the stent 10 against the interior surface of the lumen 100 and/or lesion 102. The balloon 80 is preferably a high pressure balloon (operating at 12-14 Bars) and preferably has an inflated diameter that is less than or equal to the interior diameter of the stent 10 as expanded.

30 Another feature of one preferred rapid-exchange delivery system according to the present invention is in the routing of the guidewire 104 out of the inner tube 60 and cover sheath 70 at a point between the distal end 61 of the inner tube 60 and

the distal end of the support tube 54. Turning to Figure 17, a portion of a rapid-exchange delivery system proximal from the distal end 61 of the inner tube 60 is depicted which includes the cover sheath 70 and the inner tube 60 located within the lumen 72 of the cover sheath 70. The guidewire lumen 64 of the inner tube 60 terminates in a first guidewire opening 63 in the depicted embodiment. A second guidewire opening 73 is provided in the cover sheath 70.

A guide element 130 is preferably provided that extends through the second guidewire opening 73 and the first guidewire opening 63 and into the guidewire lumen 64 of the inner tube 60. As such, advancement of the proximal end 106 of the guidewire 104 towards the proximal end of the delivery system through the guidewire lumen 64 (to the left in Figure 17) moves the proximal end 106 of the guidewire 104 into a lumen 132 in the guide element 130.

It is preferred that only a portion of the guidewire 104 fit within the lumen 132 in the guide element 130. As a result, continued advancement of the guidewire 104 towards the proximal end of the delivery system forces the guide element 130 out of the first and second guidewire openings 63/73 as well as guides the proximal end 106 of the guidewire 104 through those openings. After the proximal end 106 of the guidewire 104 is threaded through the openings 63/73 in the inner tube 60 and cover sheath 70, the distal portion of the inner tube 60 and cover sheath 70 in which the guidewire 104 is contained can be advanced through the guide catheter 90 along the guidewire 104.

Although Figure 17 illustrates one embodiment of a rapid-exchange delivery system, it will be understood that the stents according to the present invention can be delivered by any delivery system, e.g., an over-the-wire delivery system or by any other suitable delivery system. Furthermore, it will also be understood that the distal portions of the delivery system as depicted in Figures 15 and 16 could be used in connection with any suitable delivery system, including, e.g., rapid-exchange or over-the-wire delivery systems.

Furthermore, the preceding specific embodiments are illustrative of the practice of the invention. It is to be understood, however, that other expedients known to those skilled in the art or disclosed herein, may be employed without departing from the scope of the appended claims.

What is claimed is:

1. A radially expandable stent for implantation within a body lumen comprising:
 - an elongated generally tubular body defining a passageway having a longitudinal axis;
 - the body comprising a plurality of circumferential support sections arranged successively along the longitudinal axis, each of the support sections having a length along the longitudinal axis;
 - each of the circumferential support sections comprising a plurality of primary bends interconnected by struts, the primary bends being located on alternating ends of the support section around the circumference of the body, each of the struts connecting successive primary bends on opposite ends of the support section and having a midpoint generally located therebetween; and
 - at least one longitudinal member connecting adjacent support sections in the body, the longitudinal member having a first end attached proximate the midpoint of one of the struts and a second end attached proximate the midpoint of one of the struts in the adjacent support section;
 - wherein the stent is radially compressible into a compressed state in which the struts are generally aligned with the longitudinal axis and radially expandable into an expanded state in which the struts and the primary bends in each of the support sections are arranged in a zigzag pattern, and further wherein the longitudinal length of the stent in the compressed state is substantially the same as the longitudinal length of the stent in the expanded state.
2. A stent according to claim 1, wherein the body comprises first, second and third support sections, and further wherein each of the longitudinal members connecting the first support section to the second support section are circumferentially offset from each of the longitudinal members connecting the second support section to the third support section.
3. A stent according to claim 1, further comprising two or more longitudinal members connecting adjacent support sections in the body.

4. A stent according to claim 1, wherein all of the support sections are in phase with each other.
- 5 5. A stent according to claim 1, wherein at least one pair of adjacent support sections are out of phase with each other.
6. A stent according to claim 1, wherein the body comprises a nickel titanium alloy.
- 10 7. A stent according to claim 1, wherein each primary bend of the plurality of primary bends connects a pair of struts in the support section, and further wherein each pair of struts abut at a point between the primary bend and the midpoint of each of the struts in the pair of struts when the stent is in the
- 15 compressed state, whereby the bending stress is reduced at each primary bend of the plurality of primary bends.
8. A stent according to claim 7, wherein at least one of the struts in each pair of struts associated with one of the primary bends comprises a secondary
- 20 bend located between the midpoint and one end of the strut, the secondary bend including an apex facing the other strut in the pair of struts, and further wherein the point at which the pair of struts abut is at the apex of the secondary bend when the stent is in the compressed state.
- 25 9. A stent according to claim 7, wherein each strut of the plurality of struts comprises two secondary bends, one of the secondary bends located on each side of the midpoint of the strut and each of the secondary bends being spaced from the ends of the strut, each of the secondary bends having an apex, wherein the apexes of each of the secondary bends face the opposing struts in each pair of
- 30 struts associated with one of the primary bends in the support section, and further wherein the point at which each pair of struts associated with one of the primary

bends abut when the stent is in the compressed state is at the apexes of the secondary bends of the struts.

10. A stent according to claim 7, wherein at least one of the struts in each pair of struts associated with one of the primary bends comprises a protrusion located between the midpoint and one end of the strut, the protrusion facing the other strut in the pair of struts, and further wherein the point at which the pair of struts abut is at the protrusion when the stent is in the compressed state.
11. A stent according to claim 7, wherein each strut of the plurality of struts comprises two protrusions, one of the protrusions located on each side of the midpoint of the strut and each of the protrusions being spaced from the ends of the strut, wherein the protrusions face the opposing struts in each pair of struts associated with one of the primary bends in the support section, and further wherein the point at which each pair of struts abut when the stent is in the compressed state is at the protrusions in each strut in the pair of adjacent struts.
12. A radially expandable stent for implantation within a body lumen comprising:
- an elongated generally tubular body defining a passageway having a longitudinal axis, the body comprising at least one circumferential support section having a length along the longitudinal axis;
- each of the circumferential support sections comprising a plurality of primary bends interconnected by struts, the primary bends being located on alternating ends of the support section around the circumference of the body, each of the struts connecting successive primary bends on opposite ends of the support section and having a midpoint generally located therebetween;
- wherein the stent is radially compressible into a compressed state and radially expandable into an expanded state in which the struts and primary bends in each of the support sections are arranged in a zigzag pattern, and further wherein each pair of adjacent struts associated with each of the primary bends abut at a point between the primary bend and the midpoint of each strut in the

pair of adjacent struts when the stent is in the compressed state, whereby the bending stress is reduced at each primary bend of the plurality of primary bends.

13. A stent according to claim 12, wherein at least one of the struts in the pair
5 of struts associated with each of the primary bends comprises a secondary bend located between the midpoint and one end of the strut, the secondary bend including an apex facing the other strut in the pair of struts, and further wherein the point at which the pair of struts abut is at the apex of the secondary bend when the stent is in the compressed state.

10

14. A stent according to claim 12, wherein each strut of the plurality of struts comprises two secondary bends, one of the secondary bends located on each side of the midpoint of the strut and each of the secondary bends being spaced from the ends of the strut, each of the secondary bends having an apex, wherein the
15 apexes of each of the secondary bends face the opposing struts in each pair of struts associated with one of the primary bends in the support section, and further wherein the point at which each pair of struts associated with one of the primary bends abut when the stent is in the compressed state is at the apexes of the secondary bends of the struts.

20

15. A stent according to claim 12, wherein at least one of the struts in each pair of struts associated with one of the primary bends comprises a protrusion located between the midpoint and one end of the strut, the protrusion facing the other strut in the pair of struts, and further wherein the point at which the pair of
25 struts abut is at the protrusion when the stent is in the compressed state.

16. A stent according to claim 12, wherein each strut of the plurality of struts comprises two protrusions, one of the protrusions located on each side of the midpoint of the strut and each of the protrusions being spaced from the ends of the strut, wherein the protrusions face the opposing struts in each pair of struts
30 associated with one of the primary bends in the support section, and further wherein the point at which each pair of struts associated with one of the primary

bends abut when the stent is in the compressed state is at the protrusions in the struts.

17. A radially expandable stent for implantation within a body lumen
5 comprising:

an elongated generally tubular body defining a passageway having a longitudinal axis, the body comprising at least one circumferential support section having a length along the longitudinal axis;

- each of the circumferential support sections comprising a substantially
10 continuous element including a plurality of primary bends interconnected by struts, the primary bends being located on alternating ends of the support section around the circumference of the body, each of the struts connecting successive primary bends on opposite ends of the support section and having a midpoint generally located therebetween, wherein the stent is radially compressible into a
15 compressed state and radially expandable into an expanded state in which the struts and primary bends in each of the support sections are arranged in a zigzag pattern; and

means for reducing bending stress at the primary bends when the stent is in the compressed state.

20

18. A stent according to claim 17, wherein the means for reducing bending stress causes each pair of struts associated with one of the primary bends to abut at a point between the midpoints of the pair of struts and the primary bend.

- 25 19. A stent according to claim 17, wherein the body comprises a plurality of circumferential support sections arranged successively along the longitudinal axis, and further wherein the body comprises at least one longitudinal member connecting adjacent support sections in the body, the longitudinal member having a first end attached to one of the support sections and a second end
30 attached to the adjacent support section.

20. A stent according to claim 19, wherein the first end of each of the longitudinal members is attached proximate the midpoint of one of the struts in one of the support sections and the second end of each of the longitudinal members is attached proximate the midpoint of one of the struts in an adjacent support section; wherein the longitudinal length of the stent in the compressed state is substantially the same as the longitudinal length of the stent in the expanded state.
21. A delivery system for implantation of a radially-expandable stent within a body lumen comprising:
- an inner tube having a proximal end and a distal end, the inner tube having an inner tube lumen formed therein, the inner tube lumen having an opening at the distal end of the inner tube;
 - a cover sheath having a proximal end and a distal end, the cover sheath comprising a wall defining a cover sheath lumen, the inner tube located within the cover sheath lumen;
 - a stent positioned about the inner tube at the distal end of the cover sheath;
 - a first guidewire opening in the inner tube lumen, the first guidewire opening spaced from the distal end of the inner tube;
 - a second guidewire opening in the wall of the cover sheath, the second guidewire opening located proximate the first guidewire opening; and
 - a guide element having a distal end located within the inner tube lumen, the guide element extending between the first and second guidewire openings.
22. A system according to claim 21, wherein the guide element comprises a proximal end located outside of the cover sheath lumen such that the guide element extends through the second guidewire opening.
23. A system according to claim 21, wherein the guide element is removably positioned in the inner tube lumen and the first and second guidewire openings.

24. A system according to claim 21, wherein the inner tube lumen terminates at the first guidewire opening.
25. A system according to claim 21, wherein the first guidewire opening is
5 formed in an inner tube wall.
26. A system according to claim 21, wherein the guide element comprises a guide lumen formed in the distal end of the guide element.
- 10 27. A system according to claim 26, wherein the guide lumen can receive only a portion of the guidewire.
28. A system according to claim 21, wherein the stent is self-expanding and comprises:
- 15 an elongated generally tubular body defining a passageway having a longitudinal axis, the body comprising at least one circumferential support section having a length along the longitudinal axis;
- each of the circumferential support sections comprising a plurality of primary bends interconnected by struts, the primary bends being located on
- 20 alternating ends of the support section around the circumference of the body, each of the struts connecting successive primary bends on opposite ends of the support section and having a midpoint generally located therebetween;
- wherein the stent is radially compressible into a compressed state and radially expandable into an expanded state in which the struts and primary bends
- 25 in each of the support sections are arranged in a zigzag pattern, and further wherein each pair of adjacent struts associated with each of the primary bends abut at a point between the primary bend and the midpoint of each strut in the pair of adjacent struts when the stent is in the compressed state, whereby the bending stress is reduced at each primary bend of the plurality of primary bends.
- 30 29. A system according to claim 28, wherein the stent is biased against the interior surface of the cover sheath in the compressed state, and further wherein

the inner tube comprises a shoulder having an outside diameter greater than an inside diameter of the stent in the compressed state within the cover sheath, whereby movement of the cover sheath proximally forces the stent out of the cover sheath.

5

30. A system according to claim 28, wherein at least one of the struts in the pair of struts associated with each of the primary bends comprises a secondary bend located between the midpoint and one end of the strut, the secondary bend including an apex facing the other strut in the pair of struts, and further wherein
10 the point at which the pair of struts abut is at the apex of the secondary bend when the stent is in the compressed state.

31. A system according to claim 28, wherein at least one of the struts in each pair of struts associated with one of the primary bends comprises a protrusion
15 located between the midpoint and one end of the strut, the protrusion facing the other strut in the pair of struts, and further wherein the point at which the pair of struts abut is at the protrusion when the stent is in the compressed state.

32. A system according to claim 28, wherein the body comprises a plurality
20 of circumferential support sections arranged successively along the longitudinal axis, and further wherein the body comprises at least one longitudinal member connecting adjacent support sections in the body, the longitudinal member having a first end attached proximate the midpoint of one of the struts in one of the support sections and a second end attached proximate the midpoint of one of
25 the struts in an adjacent support section; wherein the longitudinal length of the stent in the compressed state is substantially the same as the longitudinal length of the stent in the expanded state.

33. A system according to claim 32, wherein the body of the stent comprises
30 first, second and third support sections, and further wherein each of the longitudinal members connecting the first support section to the second support

section are circumferentially offset from each of the longitudinal members connecting the second support section to the third support section.

34. A system according to claim 21, further comprising:
5 an expandable balloon located on the inner tube; and
an inflation lumen in fluid communication with the balloon, the inflation lumen extending from the proximal end of the delivery system to the balloon.
35. A system according to claim 34, wherein the balloon is located between
10 the stent and the inner tube.
36. A radially expandable stent for implantation within a body lumen comprising:
an elongated generally tubular body defining a passageway having a
15 longitudinal axis;
the body comprising a plurality of circumferential support sections arranged successively along the longitudinal axis, each of the support sections having a longitudinal length along the longitudinal axis;
each of the circumferential support sections comprising a plurality of
20 primary bends interconnected by struts, the primary bends being located on alternating ends of the support section around the circumference of the body, each of the struts connecting successive primary bends on opposite ends of the support section; and
at least one longitudinal member connecting adjacent support sections in
25 the body;
wherein each of the circumferential support sections is radially compressible into a compressed diameter and radially expandable into an unrestrained expanded diameter in which the struts and the primary bends in each of the support sections are arranged in a zigzag pattern, and further wherein
30 the longitudinal lengths of at least two of the circumferential support sections are different.

37. A stent according to claim 36, wherein the longitudinal lengths of successive circumferential support sections increases from one end of the stent to an opposite end of the stent.
- 5 38. A stent according to claim 37, wherein the increases in longitudinal lengths between successive support sections are uniform.
39. A stent according to claim 36, wherein the longitudinal lengths of the circumferential support sections at each end of the stent are greater than the
10 length of one of the circumferential support sections located between the circumferential support sections at each end of the stent.
40. A stent according to claim 36, wherein the circumferential support section with the shortest longitudinal length is located proximate a center of the
15 stent.
41. A stent according to claim 36, wherein the body comprises a nickel titanium alloy.
- 20 42. A radially expandable stent for implantation within a body lumen comprising:
an elongated generally tubular body defining a passageway having a longitudinal axis;
the body comprising a plurality of circumferential support sections
25 arranged successively along the longitudinal axis, each of the support sections having a longitudinal length along the longitudinal axis;
each of the circumferential support sections comprising a plurality of primary bends interconnected by struts, the primary bends being located on alternating ends of the support section around the circumference of the body,
30 each of the struts connecting successive primary bends on opposite ends of the support section; and

at least one longitudinal member connecting adjacent support sections in the body;

wherein each of the circumferential support sections is radially compressible into a compressed diameter and radially expandable into an unrestrained expanded diameter in which the struts and the primary bends in each of the support sections are arranged in a zigzag pattern, and further wherein the unrestrained expanded diameters of at least two of the support sections vary.

43. A stent according to claim 42, wherein the unrestrained expanded diameters of successive circumferential support sections increases from one end of the stent to an opposite end of the stent.

44. A stent according to claim 43, wherein the increases in the unrestrained expanded diameters between successive support sections are uniform.

45. A stent according to claim 42, wherein the unrestrained expanded diameters of the circumferential support sections at each end of the stent are smaller than the unrestrained expanded diameter of one of the circumferential support sections located between the circumferential support sections at each end of the stent.

46. A stent according to claim 42, wherein the circumferential support section with the largest unrestrained expanded diameter is located proximate a center of the stent.

47. A stent according to claim 42, wherein the longitudinal lengths of the circumferential support sections are substantially uniform.

48. A stent according to claim 42, wherein the longitudinal lengths of at least two of the circumferential support sections are different.

49. A stent according to claim 42, wherein the body comprises a nickel titanium alloy.

50. A radially expandable stent for implantation within a body lumen
5 comprising:
an elongated generally tubular body defining a passageway having a longitudinal axis;
the body comprising a plurality of circumferential support sections arranged successively along the longitudinal axis, each of the support sections
10 having a longitudinal length along the longitudinal axis;
each of the circumferential support sections comprising a plurality of primary bends interconnected by struts having a circumferential width, the primary bends being located on alternating ends of the support section around the circumference of the body, each of the struts connecting successive primary
15 bends on opposite ends of the support section; and
at least one longitudinal member connecting adjacent support sections in the body;
wherein each of the circumferential support sections is radially compressible into a compressed diameter and radially expandable into an
20 unrestrained expanded diameter in which the struts and the primary bends in each of the support sections are arranged in a zigzag pattern, and further wherein the circumferential widths of the struts in at least two of the circumferential support sections are different.

- 25 51. A stent according to claim 50, wherein the longitudinal lengths of at least two of the circumferential support sections are different.

52. A stent according to claim 50, wherein the unrestrained expanded diameters of at least two of the support sections vary.

30

53. A method of deploying a stent within a body lumen comprising:

a) providing a radially expandable stent in a compressed diameter proximate a distal end of a delivery device, wherein the stent comprises:

5 an elongated generally tubular body defining a passageway having a longitudinal axis;

the body comprising a plurality of circumferential support sections arranged successively along the longitudinal axis, each of the support sections having a longitudinal length along the longitudinal axis;

10 each of the circumferential support sections comprising a plurality of primary bends interconnected by struts, the primary bends being located on alternating ends of the support section around the circumference of the body, each of the struts connecting successive primary bends on opposite ends of the support section; and

15 at least one longitudinal member connecting adjacent support sections in the body;

wherein each of the circumferential support sections is radially compressible into the compressed diameter and radially expandable into an unrestrained expanded diameter in which the struts and the primary bends in each of the support sections are arranged in a zigzag pattern, and further wherein the longitudinal lengths of at least two of the circumferential support sections are different;

20 b) advancing the distal end of the delivery device and the stent through a body lumen; and

c) deploying the stent at a desired location within the body lumen.

54. A method of deploying a stent within a body lumen comprising:

30 a) providing a radially expandable stent in a compressed diameter proximate a distal end of a delivery device, wherein the stent comprises:

an elongated generally tubular body defining a passageway having a longitudinal axis;

the body comprising a plurality of circumferential support sections arranged successively along the longitudinal axis, each of the support sections having a longitudinal length along the longitudinal axis;

each of the circumferential support sections comprising a plurality of primary bends interconnected by struts, the primary bends being located on alternating ends of the support section around the circumference of the body, each of the struts connecting successive primary bends on opposite ends of the support section; and

at least one longitudinal member connecting adjacent support sections in the body;

wherein each of the circumferential support sections is radially compressible into the compressed diameter in which the struts are generally aligned with the longitudinal axis and radially expandable into an unrestrained expanded diameter in which the struts and the primary bends in each of the support sections are arranged in a zigzag pattern, and further wherein the unrestrained expanded diameters of at least two of the support sections vary;

b) advancing the distal end of the delivery device and the stent through a body lumen; and

c) deploying the stent at a desired location within the body lumen.

55. A method of deploying a stent within a body lumen comprising:

a) providing a radially expandable stent in a compressed diameter proximate a distal end of a delivery device, wherein the stent comprises:

an elongated generally tubular body defining a passageway having a longitudinal axis;

the body comprising a plurality of circumferential support sections arranged successively along the longitudinal axis, each of

the support sections having a longitudinal length along the longitudinal axis;

5 each of the circumferential support sections comprising a plurality of primary bends interconnected by struts, the primary bends being located on alternating ends of the support section around the circumference of the body, each of the struts connecting successive primary bends on opposite ends of the support section; and

10 at least one longitudinal member connecting adjacent support sections in the body;

wherein each of the circumferential support sections is radially compressible into the compressed diameter and radially expandable into an unrestrained expanded diameter in which the struts and the primary bends in each of the support sections are arranged in a zigzag pattern, and further wherein the circumferential width of the struts in at least two of the circumferential support sections are different;

- 15
- b) advancing the distal end of the delivery device and the stent through a body lumen; and
- 20 c) deploying the stent at a desired location within the body lumen.

56. A method of deploying a stent within a body lumen comprising:

- a) providing a radially expandable stent on a delivery system comprising:
- 25 an inner tube having a proximal end and a distal end, the inner tube having an inner tube lumen formed therein, the inner tube lumen having an opening at the distal end of the inner tube and a first guidewire opening in the inner tube lumen, the first guidewire opening spaced from the distal end of the inner tube;
- 30 a stent positioned on the exterior surface of the inner tube at the distal end of the inner tube;
- a cover sheath having a proximal end and a distal end, the cover sheath comprising a wall defining a cover sheath lumen, the inner tube

and stent located within the cover sheath lumen, the cover sheath further including a second guidewire opening in the wall of the cover sheath, the second guidewire opening located proximate the first guidewire opening in the inner tube; and

5 a guide element having a distal end located within the inner tube lumen, the guide element extending between the first and second guidewire openings, wherein the guide element comprises a guide lumen formed in the distal end of the guide element;

10 b) positioning a guidewire within a body lumen, wherein a proximal end of the guidewire extends out of the body lumen;

 c) inserting the proximal end of the guidewire into the inner tube lumen at the distal end of the inner tube;

 d) advancing the proximal end of the guidewire through the inner tube
15 lumen towards the first guidewire opening and the distal end of the guide element, wherein at least a portion of the proximal end of the guidewire is advanced into the guide lumen in the distal end of the guide element;

 e) advancing the proximal end of the guidewire through the first and second guidewire openings;

20 f) advancing the distal end of the inner tube and the stent over the guidewire towards the distal end of the guidewire, wherein the stent is positioned at a desired location within the body lumen; and

 g) deploying the stent at the desired location within the body lumen.

25 57. A method according to claim 56, wherein the guide element is removably positioned in the inner tube lumen and the first and second guidewire openings, and further wherein the step of advancing the proximal end of the guidewire through the first and second guidewire openings further comprises moving the guide element out of the first and second guidewire openings.

30

58. A method according to claim 56, wherein the guide lumen can receive only a portion of the proximal end of the guidewire.

59. A method according to claim 56, wherein the guide element comprises a proximal end located outside of the cover sheath lumen such that the guide element extends through the second guidewire opening.

5

60. A method of deploying a stent within a body lumen comprising:

a) providing a radially expandable stent on a delivery system comprising:

an inner tube having a proximal end and a distal end, the inner tube having an inner tube lumen formed therein;

10. a stent positioned on the exterior surface of the inner tube at the distal end of the inner tube;

an expandable balloon located on the inner tube;

an inflation lumen in fluid communication with the balloon, the inflation lumen extending from the proximal end of the delivery system to the balloon; and

15

a cover sheath having a proximal end and a distal end, the cover sheath comprising a wall defining a cover sheath lumen, the inner tube, stent, and balloon located within the cover sheath lumen;

b) positioning the inner tube, stent, balloon and cover sheath within a body lumen;

20

c) moving the cover sheath proximally relative to the distal end of the inner tube to deploy the stent with the body lumen; and

d) inflating the balloon within the stent.

25 61. A method according to claim 60, wherein the balloon is located between the stent and the inner tube.

62. A method of deploying a stent within a body lumen comprising:

a) providing a radially expandable stent on a delivery system comprising:

30 an inner tube having a proximal end and a distal end;

a stent positioned on the exterior surface of the inner tube at the distal end of the inner tube;

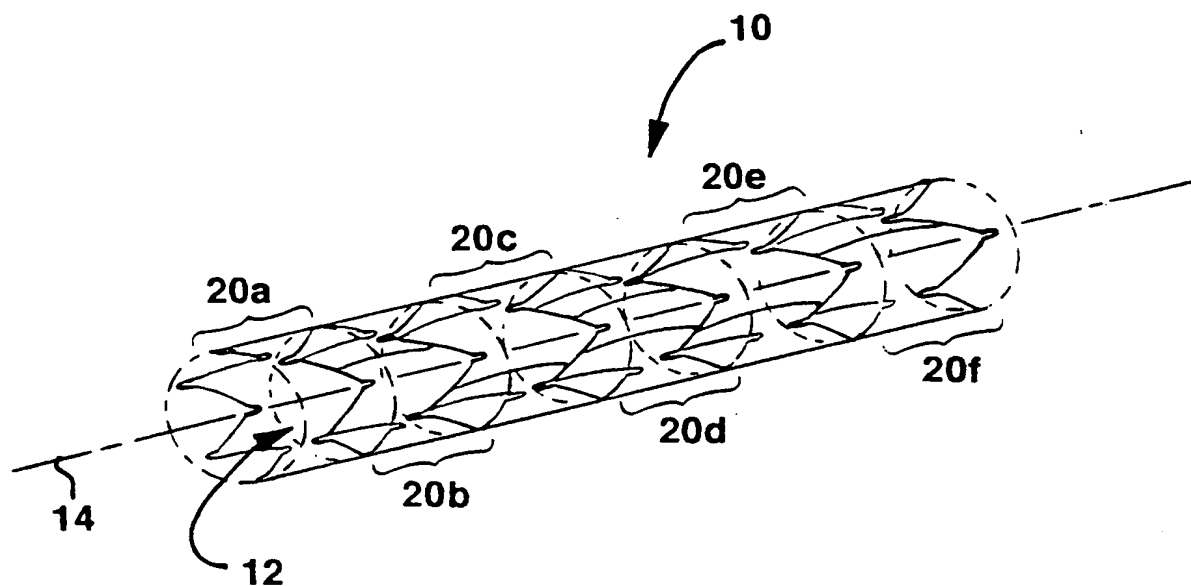
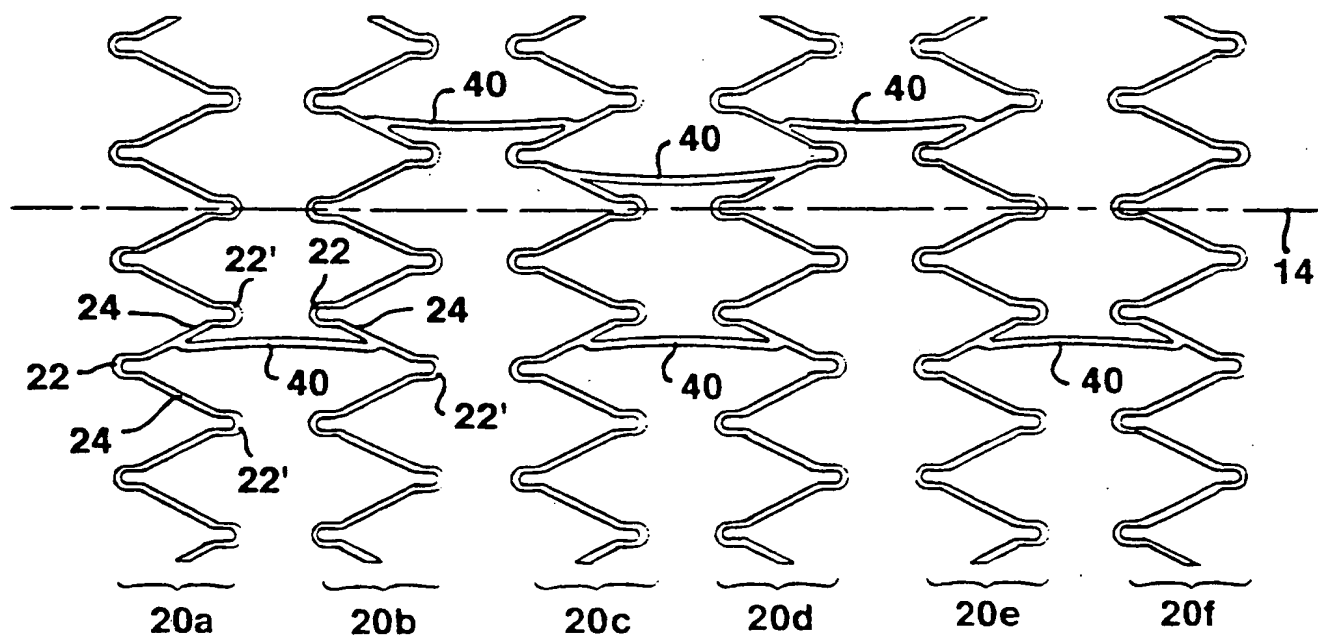
a cover sheath having a proximal end and a distal end, the cover sheath including a cover sheath lumen, the inner tube and stent located within the cover sheath lumen; and

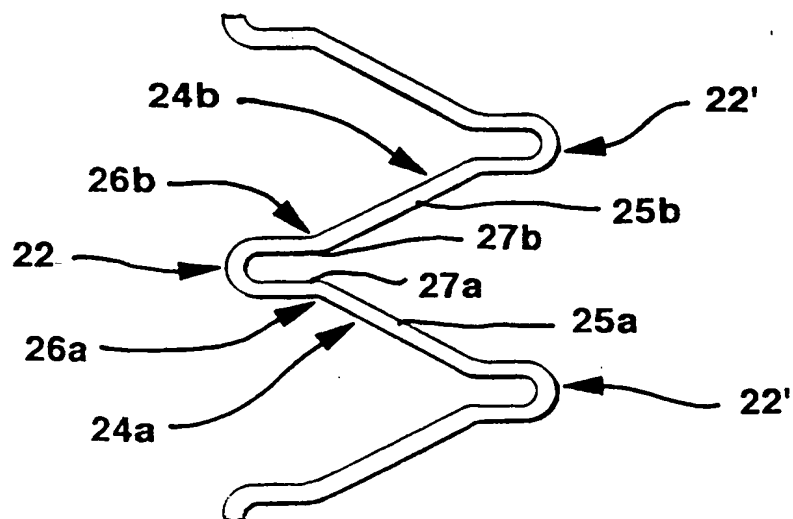
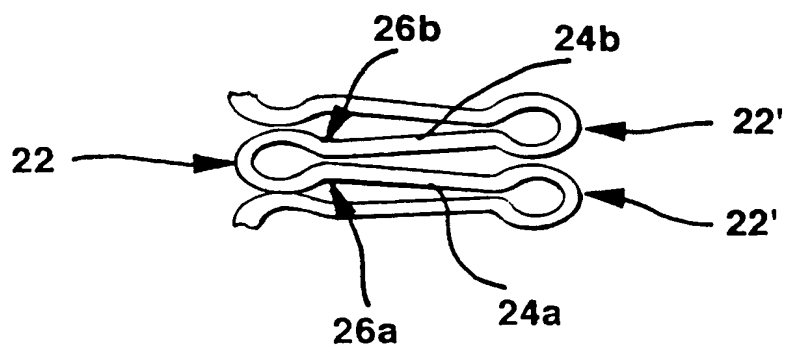
5 a support tube having a proximal end and a distal end, the support tube including a support tube lumen containing at least a portion of the proximal end of the cover sheath, the cover sheath being movable in the proximal and distal directions within the support tube lumen and the position of the inner tube being fixed relative to the position of the support tube;

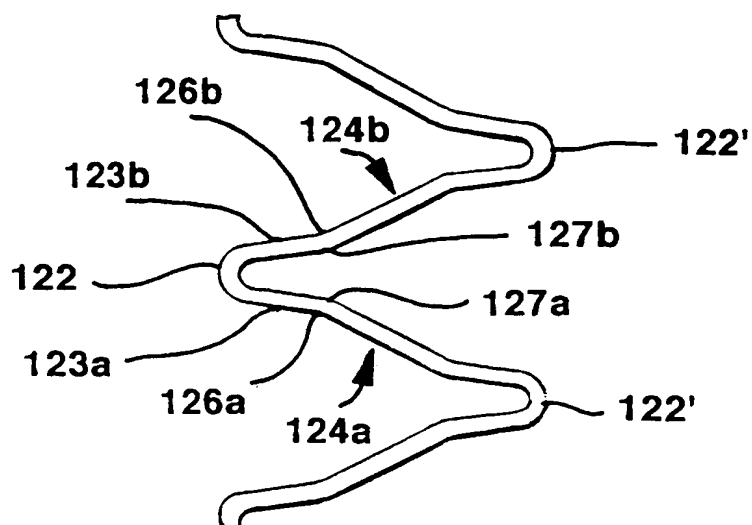
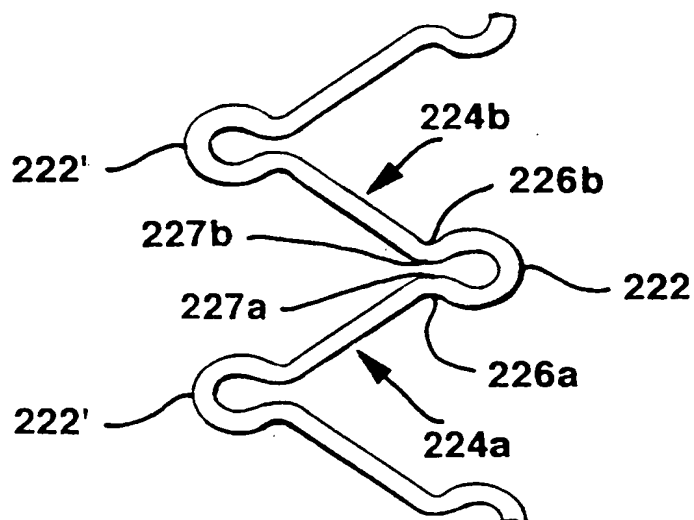
- 10 b) positioning a guide catheter within a body lumen;
c) advancing the distal ends of the inner tube and the cover sheath through the guide catheter;
d) fixing the position of the support tube relative to the guide catheter, wherein the positions of the distal end of the inner tube and the stent within the
15 body lumen are also fixed relative to the guide catheter; and
e) moving the cover sheath proximally to release the stent on the distal end of the inner tube, thereby deploying the stent within the body lumen.

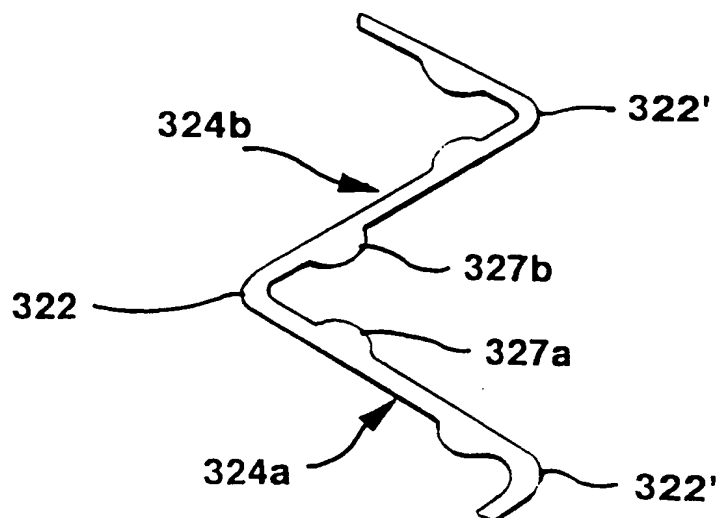
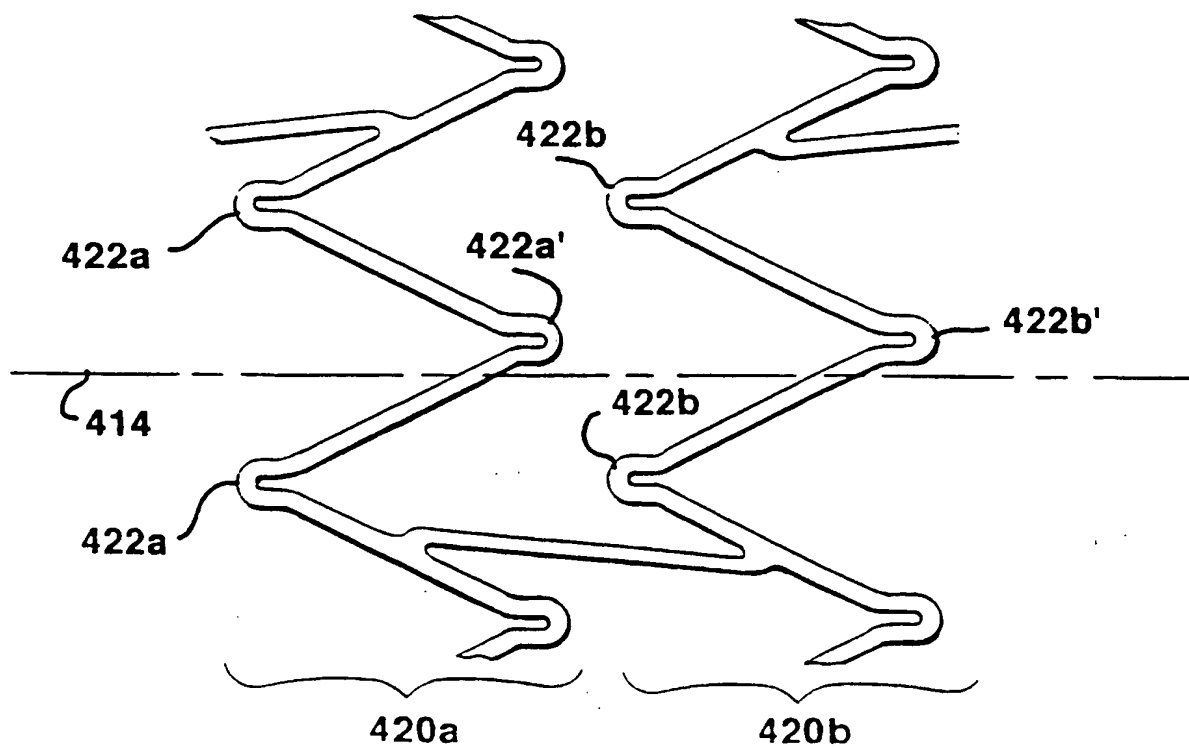
63. A method according to claim 62, wherein the guide catheter includes a
20 hemostasis valve and further wherein the step of fixing the position of the support tube relative to the guide catheter comprises closing the hemostasis valve on the support tube.

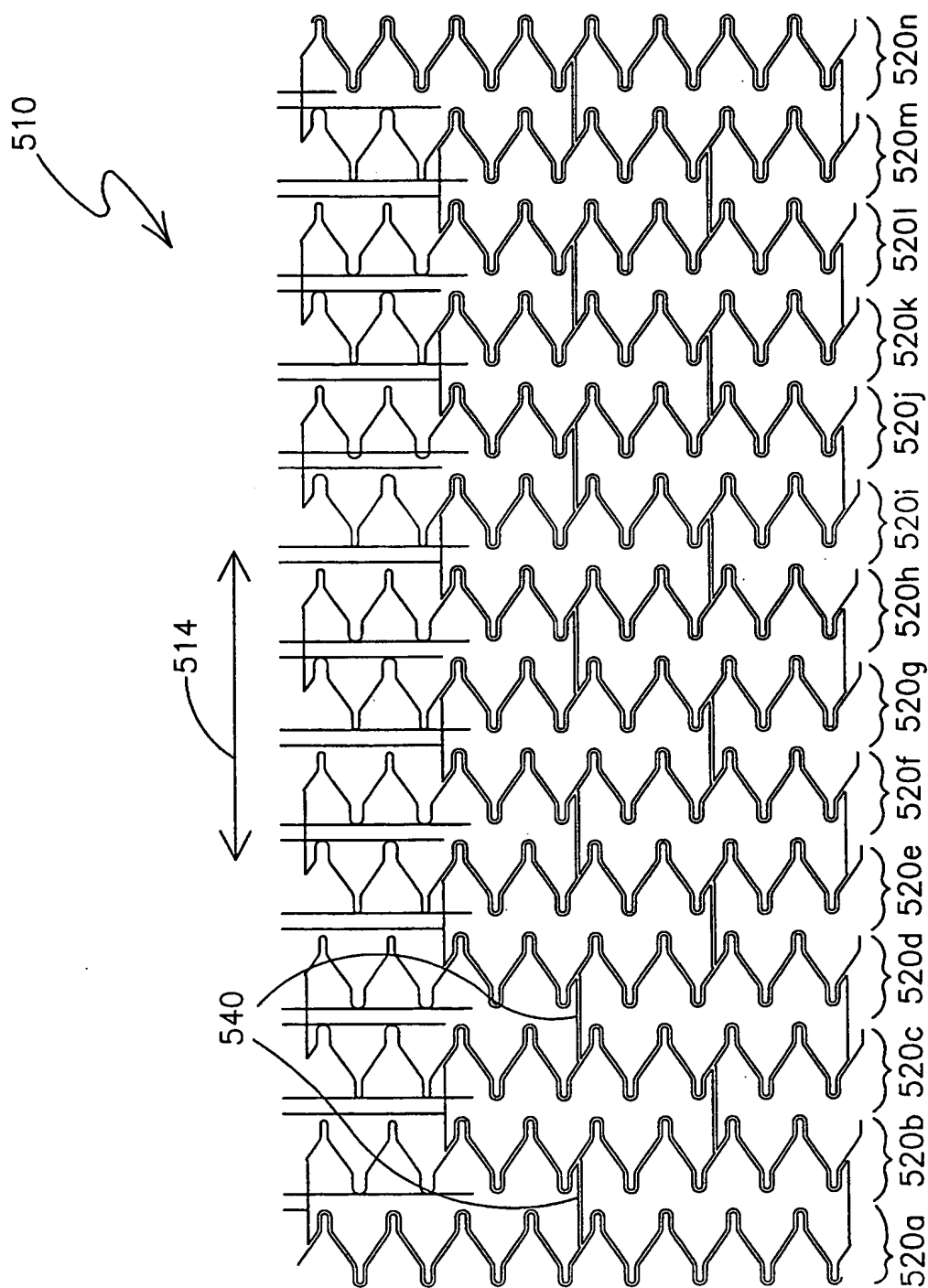
64. A method according to claim 62, wherein the support tube and the inner
25 tube are fixedly attached to a handle and the cover sheath is attached to an actuator, the actuator movable relative to the handle, and further wherein the step of moving the cover sheath comprises moving the actuator relative to the handle.

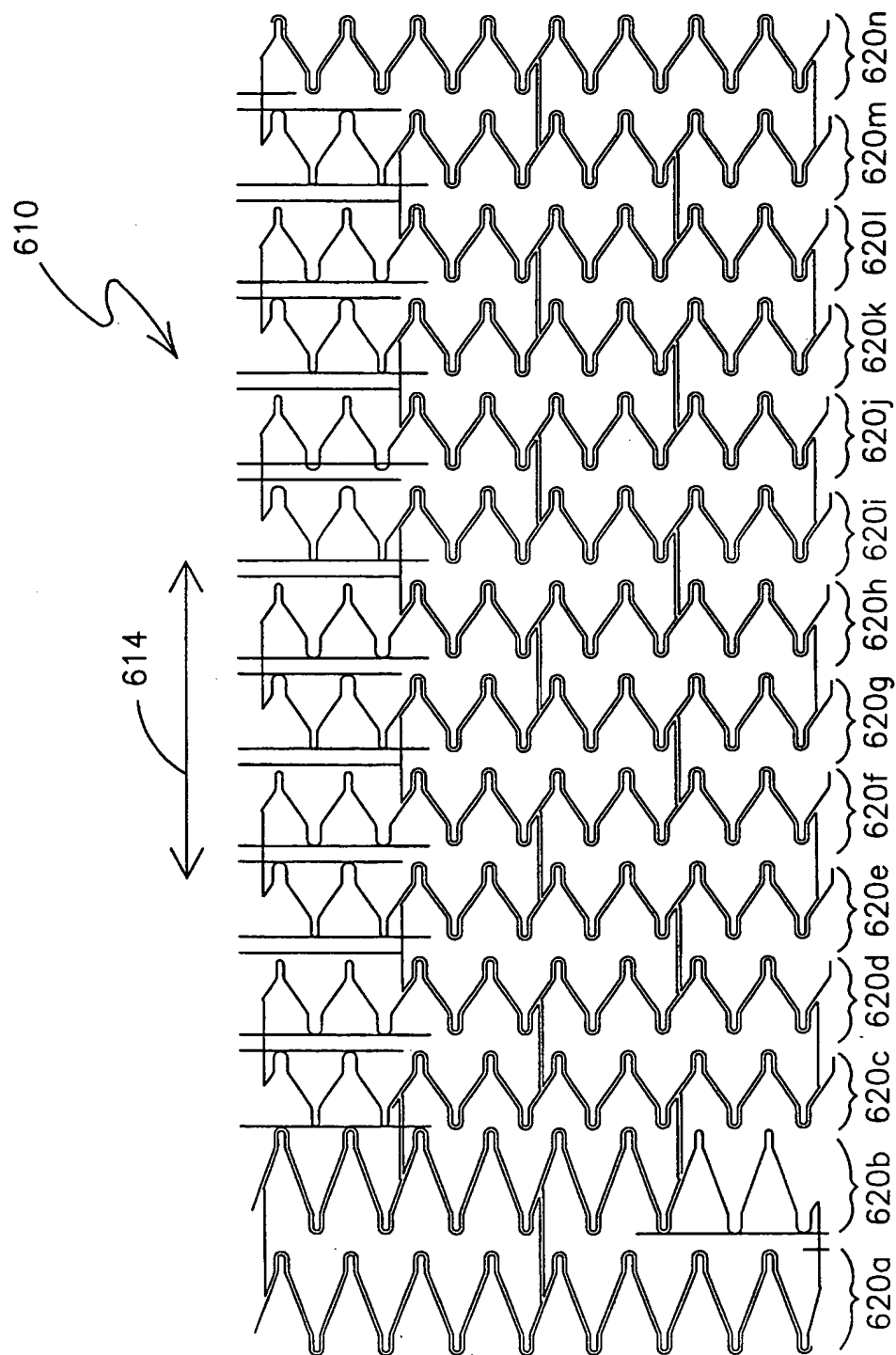
**FIG. 1****FIG. 2**

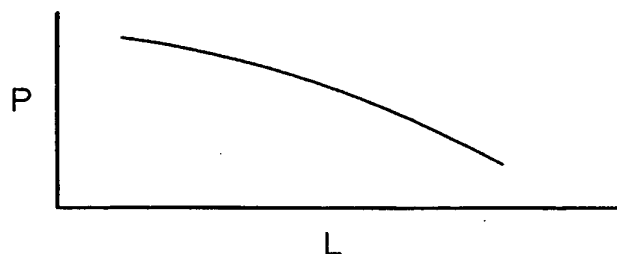
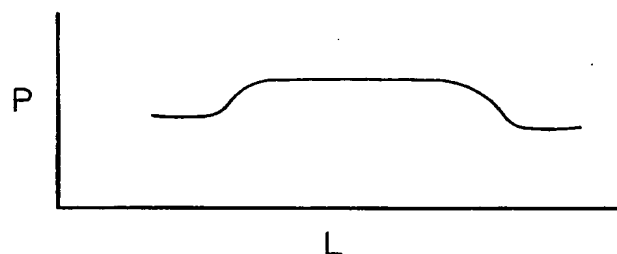
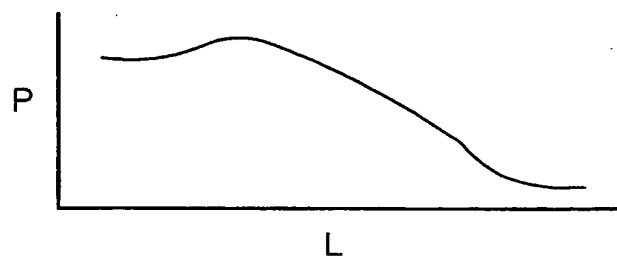
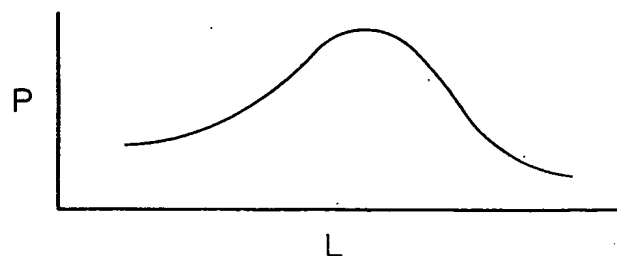
**FIG. 3****FIG. 4**

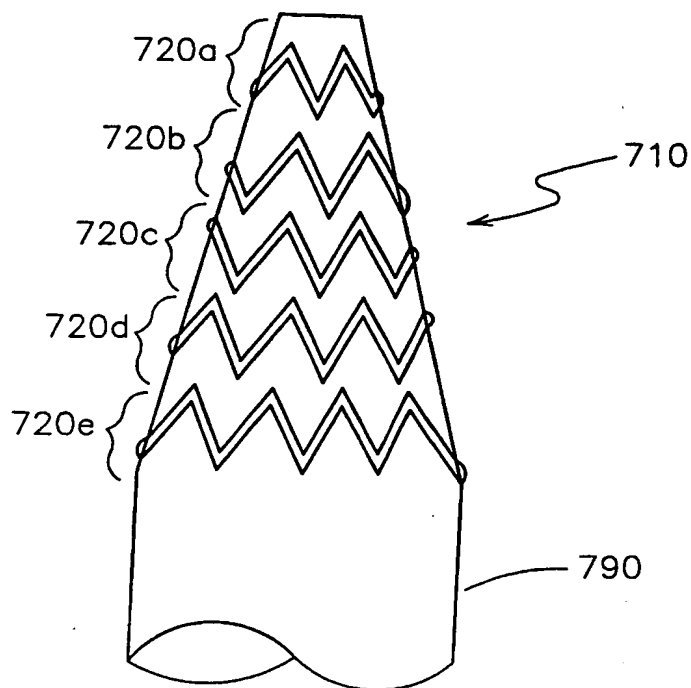
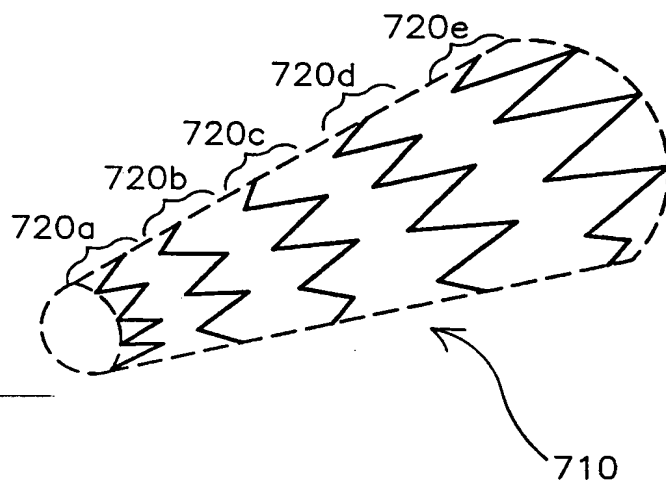
**FIG. 5****FIG. 6**

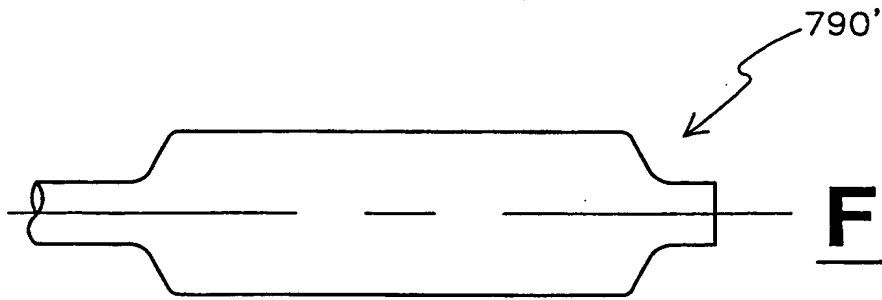
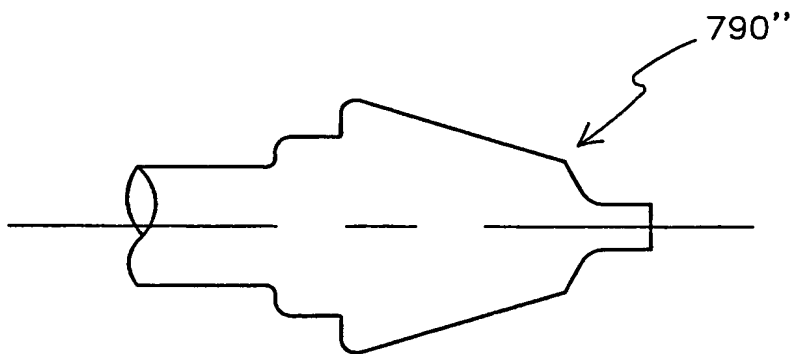
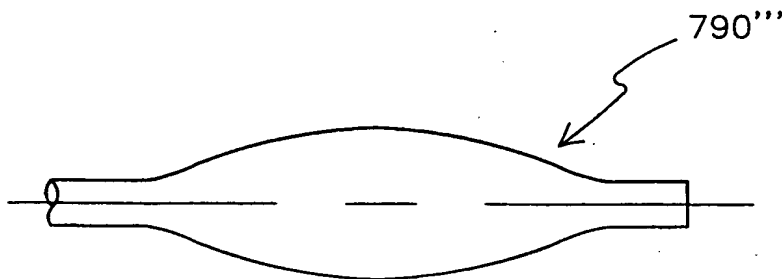
**FIG. 7****FIG. 8**

**FIG. 9**

**FIG. 10**

**FIG. 11A****FIG. 11B****FIG. 11C****FIG. 11D**



**FIG. 12B****FIG. 12C****FIG. 12D**

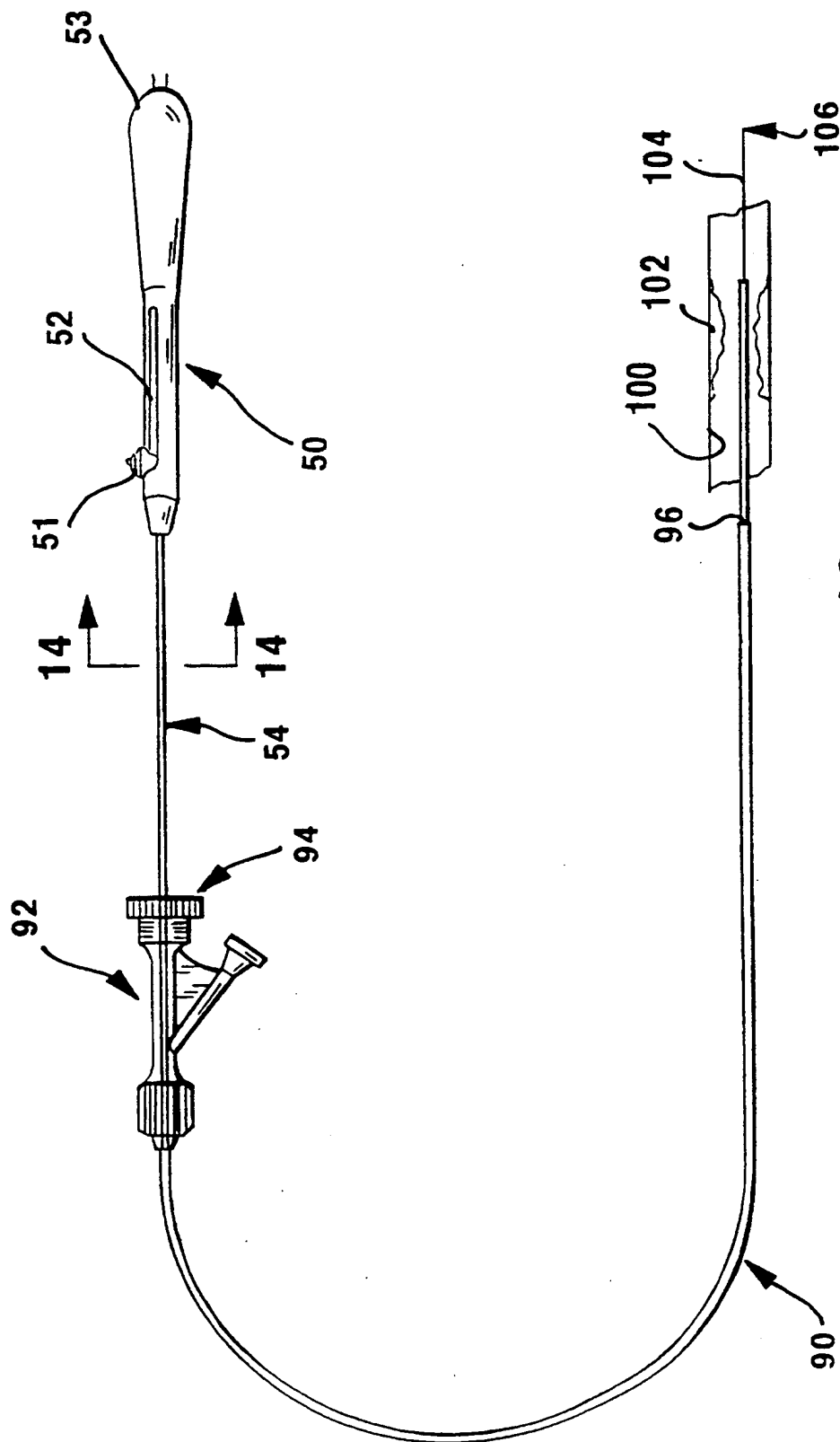


FIG. 13

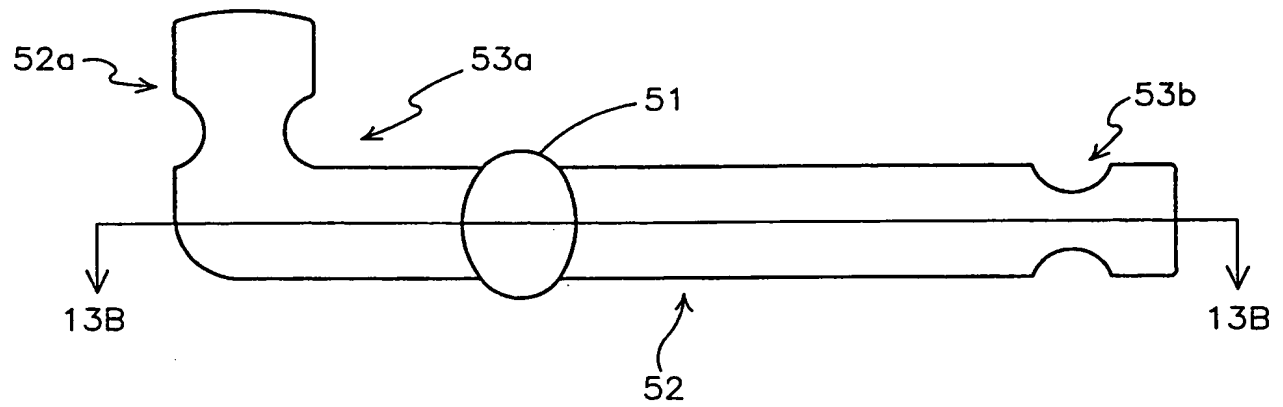


FIG. 13A

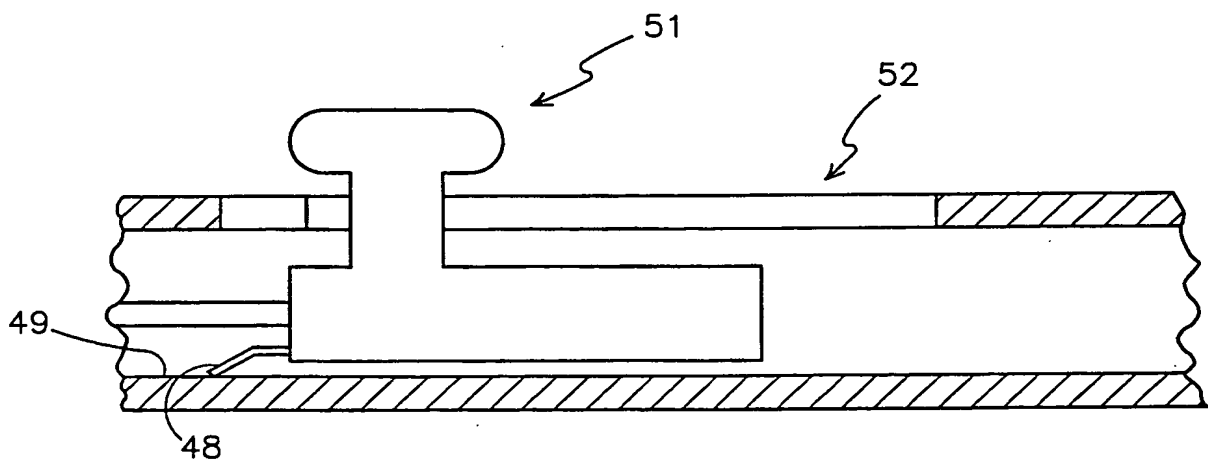


FIG. 13B

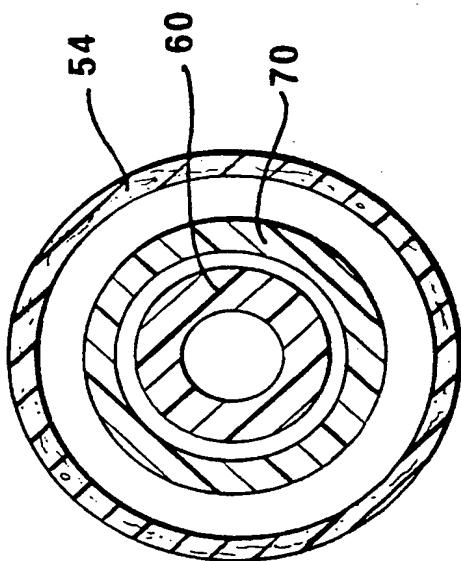


FIG. 14

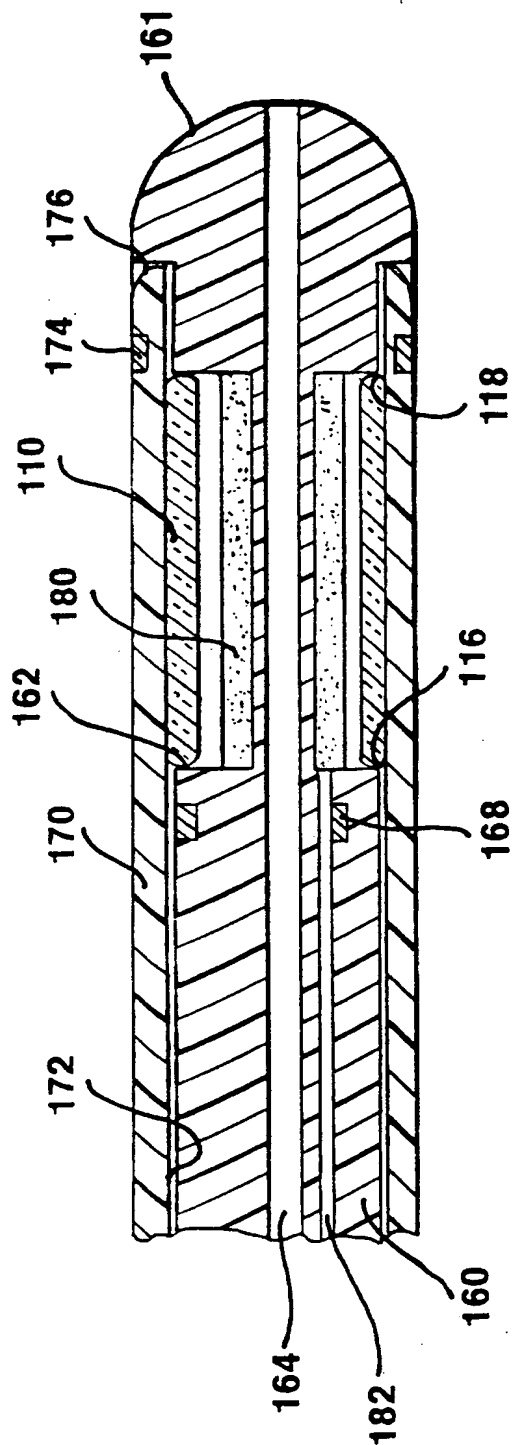
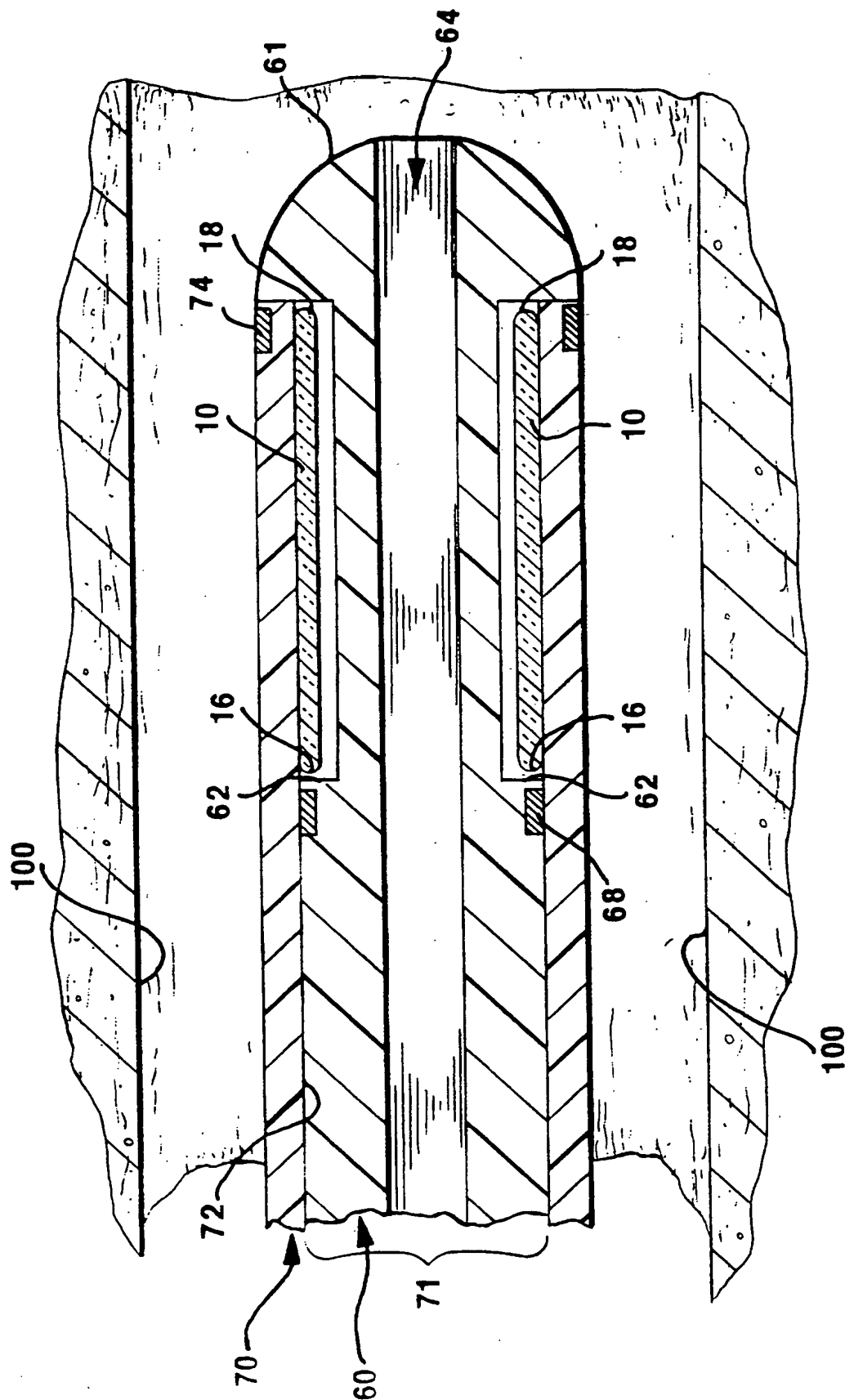


FIG. 16



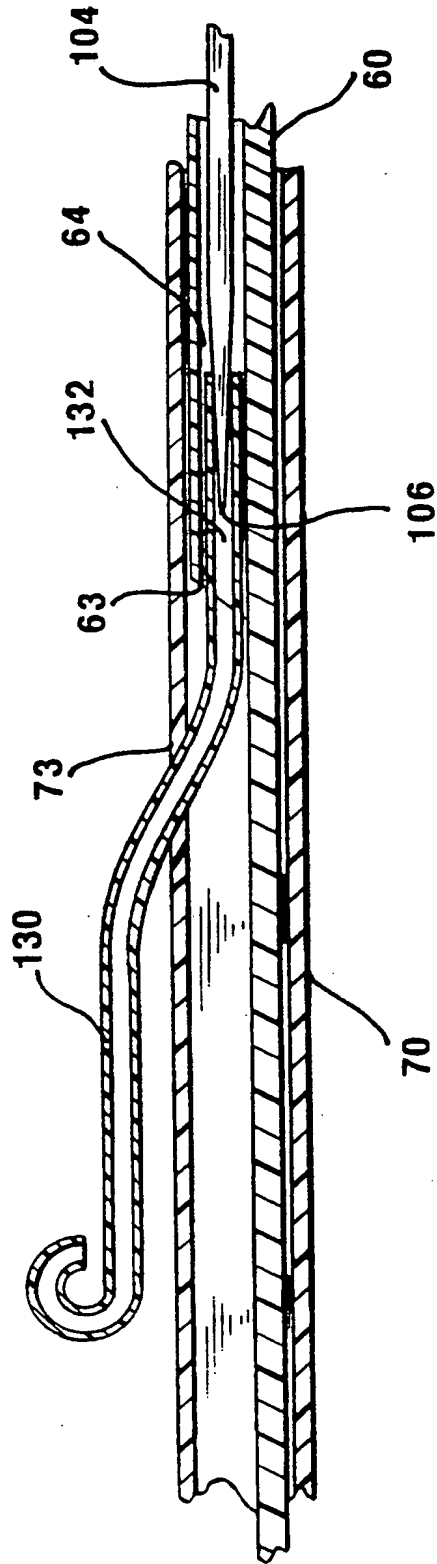


FIG.17

1
INTERNATIONAL SEARCH REPORT

International application No.

PCT/US 99/02453

A. CLASSIFICATION OF SUBJECT MATTER

IPC6: A61F 2/06 // A61M 25/00

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC6: A61F

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

EPODOC

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	EP 0818184 A1 (B. BRAUN CELSA), 14 January 1998 (14.01.98), column 3, line 36 - column 4, line 12; column 6, line 43 - column 7, line 5, figures 1-2, 12	1-20
A	--	36-52
X	WO 9415549 A1 (SCHNEIDER (USA)INC.), 21 July 1994 (21.07.94)	21-27, 34-35
A	--	28-33



Further documents are listed in the continuation of Box C.



See patent family annex.

* Special categories of cited documents:

"A" document defining the general state of the art which is not considered to be of particular relevance

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"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

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"&" document member of the same patent family

Date of the actual completion of the international search

26 July 1999

Date of mailing of the international search report

15.09.1999

Name and mailing address of the International Searching Authority
European Patent Office P.B. 5818 Patentlaan 2
NL-2280 HV Rijswijk
Tel(+31-70)340-2040. Tx 31 651 epo nl,
Fax(+31-70)340-3016

Authorized officer

LEIF BRANDER

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US 99/02453

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages.	Relevant to claim No.
A	WO 9641592 A1 (BARD GALWAY LIMITED), 27 December 1996 (27.12.96), page 7, line 11 - page 10, line 13, figures 1-3 -- -----	1-20, 36-52

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US 99/ 02453

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.: 53-64
because they relate to subject matter not required to be searched by this Authority, namely:
Rule 39.1(iv) PCT - Method for treatment of the human or animal body by surgery or therapy
2. ☐ Claims Nos.:
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

Claims 1-20: A radially expandable stent with mid point-attached struts.
Claims 21-35: A delivery system.
Claims 36-52: A radially expandable stent with different portions.

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. ☒ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

S/ 8844

INTERNATIONAL SEARCH REPORT

Information on patent family members

01/07/99

International application No.

PCT/US 99/02453

Patent document cited in search report			Publication date	Patent family member(s)		Publication date
EP	0818184	A1	14/01/98	FR	2750853 A,B	16/01/98
				US	5897589 A	27/04/99

WO	9415549	A1	21/07/94	CA	2149887 A	21/07/94
				DE	9321003 U	10/08/95
				EP	0676936 A	18/10/95
				JP	8500757 T	30/01/96
				US	5690644 A	25/11/97

WO	9641592	A1	27/12/96	AU	5675196 A	09/01/97
				AU	5776696 A	09/01/97
				CA	2223479 A	27/12/96
				CA	2223502 A	27/12/96
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				EP	0830109 A	25/03/98
				IL	122202 D	00/00/00
				IL	122203 D	00/00/00
				IT	1283328 B	16/04/98
				IT	B0950292 A	09/12/96
				WO	9641591 A	27/12/96
				IT	1285898 B	26/06/98
				IT	B0960201 A	15/10/97
				IT	1285899 B	26/06/98
				IT	B0960202 A	15/10/97
